

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
25 July 2002 (25.07.2002)

PCT

(10) International Publication Number
WO 02/057236 A1

(51) International Patent Classification⁷: C07D 211/76,
211/74, 265/32, 241/08, 401/04, 263/22, 237/14, 223/10,
A61K 31/535, 31/50, 31/4412, 31/421

(74) Common Representative: MERCK PATENT GMBH;
Frankfurter Strasse 250, 64293 Darmstadt (DE).

(21) International Application Number: PCT/EP01/14296

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,
CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,
MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI,
SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU,
ZA, ZW.

(22) International Filing Date: 5 December 2001 (05.12.2001)

(84) Designated States (*regional*): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),
Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),
European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR,
GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent
(BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR,
NE, SN, TD, TG).

(25) Filing Language: English

Published:

(26) Publication Language: English

- with international search report
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments

(30) Priority Data:
101 02 322.7 19 January 2001 (19.01.2001) DE

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

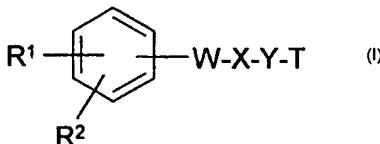
(71) Applicant (*for all designated States except US*): MERCK
PATENT GMBH [DE/DE]; Frankfurter Strasse 250,
64293 Darmstadt (DE).

(72) Inventors; and

(75) Inventors/Applicants (*for US only*): DORSCH, Dieter
[DE/DE]; Königsberger Str. 17A, 64372 Ober-Ramstadt
(DE). MEDERSKI, Werner [DE/DE]; Katzenelbogenweg 1,
64673 Zwingenberg (DE). TSAKLAKIDIS, Christos [GR/DE];
Im Langewann 54, 69469 Weinheim (DE). CEZANNE, Bertram
[DE/DE]; Goethestrasse 47, 64546 Mörfelden-Walldorf (DE).
GLEITZ, Johannes [DE/DE]; Liebigstrasse 26, 64293 Darmstadt (DE).
BARNES, Christopher [GB/DE]; Alleestrasse 21, 65812
Bad Soden (DE).

(54) Title: PHENYL DERIVATIVES

WO 02/057236 A1

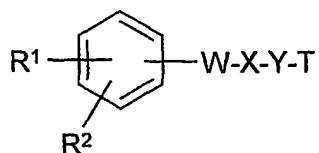


(57) Abstract: Novel compounds of the formula I in which W, X, Y, T, R¹ and R² are as defined in Patent Claim 1, are inhibitors of coagulation factor Xa and can be employed for the prophylaxis and/or therapy of thromboembolic disorders.

Phenyl derivatives

The invention relates to compounds of the formula I

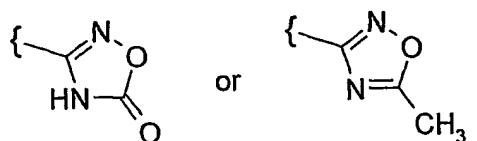
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10 in which

R^1 is CN , $CON(R^3)_2$, $-[C(R^4)_2]_nN(R^3)_2$,
 $-C(=NH)-NH_2$ which is unsubstituted or monosubstituted by
 $C(=O)R^3$, $COOR^3$, OR^3 , $OCOR^3$, $OCOOR^3$ or by a conventional
15 amino-protecting group, or is

20



30

R^2 is H , Hal , A , OR^3 , $N(R^3)_2$, NO_2 , CN , $COOR^3$, $CON(R^3)_2$,
 $-[C(R^4)_2]_n-Ar$, $-[C(R^4)_2]_n-Het$ or $-[C(R^4)_2]_n$ -cycloalkyl,
25 R^3 is H , A , $-[C(R^4)_2]_n-Ar$, $-[C(R^4)_2]_n-Het$ or $-[C(R^4)_2]_n$ -cycloalkyl,
 R^4 is H or A ,
 W is $-C(R^3)_2-$, $-[C(R^3)_2]_2-$, $-OC(R^3)_2-$ or $-NR^3C(R^3)_2-$,
 X is $CONR^3$, $CONR^3C(R^4)_2-$, $-C(R^4)_2NR^3-$, $-C(R^4)_2NR^3C(R^4)_2-$,
 $-C(R^4)_2O-$ or $-C(R^4)_2OC(R^4)_2-$,
30 Y is alkylene, cycloalkylene, Het-diyl or Ar-diyl,
 T is OR^3 , $N(R^3)_2$ or a monocyclic or bicyclic, saturated, unsaturated
or aromatic heterocyclic radical having from 1 to 4 N, O and/or S
atoms which is unsubstituted or monosubstituted, disubstituted or
trisubstituted by Hal , A , $-[C(R^4)_2]_n-Ar$, $-[C(R^4)_2]_n-Het$,

35

-[C(R⁴)₂]_n-cycloalkyl, OR³, N(R³)₂, NO₂, CN, COOR³, CON(R³)₂, NR³COA, NR³SO₂A, COR³, SO₂NR³, S(O)_mA and/or carbonyl oxygen,

- 5 A is unbranched or branched alkyl having 1-6 carbon atoms, in which one or two CH₂ groups may be replaced by O or S atoms and/or by -CH=CH- groups and/or, in addition, 1-7 H atoms may be replaced by F,
 - 10 Ar is phenyl, naphthyl or biphenyl, each of which is unsubstituted or monosubstituted, disubstituted or trisubstituted by Hal, A, OR⁴, N(R⁴)₂, NO₂, CN, COOR⁴, CON(R⁴)₂, NR⁴COA, NR⁴SO₂A, COR⁴, SO₂NR⁴, S(O)_mA, -[C(R⁴)₂]_n-COOR³ or O-[C(R⁴)₂]_o-COOR³,
 - 15 Het is a monocyclic or bicyclic, saturated, unsaturated or aromatic heterocyclic radical having from 1 to 4 N, O and/or S atoms which is unsubstituted or monosubstituted, disubstituted or trisubstituted by Hal, A, -[C(R⁴)₂]_n-Ar, -[C(R⁴)₂]_n-Het', -[C(R⁴)₂]_n-cycloalkyl, OR³, N(R³)₂, NO₂, CN, COOR³, CON(R³)₂, NR³COA, NR³SO₂A, COR³, SO₂NR³, S(O)_mA and/or carbonyl oxygen,
 - 20 Het' is a monocyclic or bicyclic, saturated, unsaturated or aromatic heterocyclic radical having from 1 to 4 N, O and/or S atoms which is unsubstituted or monosubstituted or disubstituted by Hal, A, OR³, N(R³)₂, NO₂, CN, COOR³, CON(R³)₂, NR³COA, NR³SO₂A, COR³, SO₂NR³, S(O)_mA and/or carbonyl oxygen,
 - 25 Hal is F, Cl, Br or I,
 - m and
 - 30 n are each, independently of one another, 0, 1 or 2,
 - o is 1, 2 or 3,
- and their pharmaceutically usable derivatives, solvates and stereoisomers, including mixtures thereof in all ratios.

The invention had the object of finding novel compounds having valuable properties, in particular those which can be used for the preparation of medicaments.

- 5 It has been found that the compounds of the formula I and their salts have very valuable pharmacological properties and are well tolerated. In particular, they exhibit factor Xa-inhibiting properties and can therefore be employed for combating and preventing thromboembolic disorders, such as thrombosis, myocardial infarction, arteriosclerosis, inflammation, apoplexia, angina pectoris, restenosis after angioplasty and claudicatio intermittens.
- 10 15 The compounds of the formula I according to the invention may furthermore be inhibitors of the coagulation factors factor VIIa, factor IXa and thrombin in the blood coagulation cascade.
- 20 25 Aromatic amidine derivatives having an antithrombotic action are disclosed, for example, in EP 0 540 051 B1, WO 00/71508, WO 00/71511, WO 00/71493, WO 00/71507, WO 00/71509, WO 00/71512, WO 00/71515 and WO 00/71516. Cyclic guanidines for the treatment of thromboembolic disorders are described, for example, in WO 97/08165. Aromatic heterocyclic compounds having factor Xa-inhibitory activity are disclosed, for example, in WO 96/10022. Substituted N-[(aminoimino-methyl)phenylalkyl]azaheterocycllamides as factor Xa inhibitors are described in WO 96/40679.
- 30

The antithrombotic and anticoagulant effect of the compounds according to the invention is attributed to the inhibitory action against activated coagulation protease, known by the name factor Xa, or to the inhibition of

other activated serine proteases, such as factor VIIa, factor IXa or thrombin.

5 Factor Xa is one of the proteases involved in the complex process of blood coagulation. Factor Xa catalyses the conversion of prothrombin into thrombin. Thrombin cleaves fibrinogen into fibrin monomers, which, after crosslinking, make an elementary contribution to thrombus formation.
10 Activation of thrombin may result in the occurrence of thromboembolic disorders. However, inhibition of thrombin may inhibit the fibrin formation involved in thrombus formation. The inhibition of thrombin can be measured, for example, by the method of G.F. Cousins et al. in *Circulation* 1996, 94, 1705-1712.

15

Inhibition of factor Xa can thus prevent the formation of thrombin. The compounds of the formula I according to the invention and their salts engage in the blood coagulation process by inhibiting factor Xa and thus inhibit the formation of thrombuses.

20
25 The inhibition of factor Xa by the compounds according to the invention and the measurement of the anticoagulant and antithrombotic activity can be determined by conventional in-vitro or in-vivo methods. A suitable method is described, for example, by J. Hauptmann et al. in *Thrombosis and Haemostasis* 1990, 63, 220-223.

30 The inhibition of factor Xa can be measured, for example, by the method of T. Hara et al. in *Thromb. Haemostas.* 1994, 71, 314-319.

35

Coagulation factor VIIa initiates the extrinsic part of the coagulation cascade after binding to tissue factor and contributes to the activation of

factor X to give factor Xa. Inhibition of factor VIIa thus prevents the formation of factor Xa and thus subsequent thrombin formation.

5 The inhibition of factor VIIa by the compounds according to the invention and the measurement of the anticoagulant and antithrombotic activity can be determined by conventional in-vitro or in-vivo methods. A conventional method for the measurement of the inhibition of factor VIIa is described, for example, by H. F. Ronning et al. in *Thrombosis Research* 1996, 84, 73–81.

10

Coagulation factor IXa is generated in the intrinsic coagulation cascade and is likewise involved in the activation of factor X to give factor Xa.

15 Inhibition of factor IXa can therefore prevent the formation of factor Xa in a different way.

The inhibition of factor IXa by the compounds according to the invention and the measurement of the anticoagulant and antithrombotic activity can be determined by conventional in-vitro or in-vivo methods. A suitable 20 method is described, for example, by J. Chang et al. in *Journal of Biological Chemistry* 1998, 273, 12089-12094.

25 The compounds according to the invention may furthermore be used for the treatment of tumours, tumour illnesses and/or tumour metastases. A correlation between tissue factor TF / factor VIIa and the development of various types of cancer has been indicated by T.Taniguchi and N.R. Lemoine in Biomed. Health Res. (2000), 41 (Molecular Pathogenesis of 30 Pancreatic Cancer), 57-59.

30 The publications listed below describe an antitumoural action of TF-VII and factor Xa inhibitors for various types of tumour:
K.M. Donnelly et al. in Thromb. Haemost. 1998; 79: 1041-1047;
E.G. Fischer et al. in J. Clin. Invest. 104: 1213-1221 (1999);
35 B.M. Mueller et al. in J. Clin. Invest. 101: 1372-1378 (1998);

M.E. Bromberg et al. in Thromb. Haemost. 1999; 82: 88-92

- The compounds of the formula I can be employed as medicament active ingredients in human and veterinary medicine, in particular for the treatment and prevention of thromboembolic disorders, such as thrombosis, myocardial infarction, arteriosclerosis, inflammation, apoplexia, angina pectoris, restenosis after angioplasty, claudicatio intermittens, venous thrombosis, pulmonary embolism, arterial thrombosis, myocardial ischaemia, unstable angina and strokes based on thrombosis.
- The compounds according to the invention are also employed for the treatment or prophylaxis of atherosclerotic diseases, such as coronary arterial disease, cerebral arterial disease or peripheral arterial disease.
- The compounds are also employed in combination with other thrombolytic agents in the case of myocardial infarction, furthermore for prophylaxis for reocclusion after thrombolysis, percutaneous transluminal angioplasty (PTCA) and coronary bypass operations.
- The compounds according to the invention are furthermore used for the prevention of rethrombosis in microsurgery, furthermore as anticoagulants in connection with artificial organs or in haemodialysis.
- The compounds are furthermore used in the cleaning of catheters and medical aids *in vivo* in patients, or as anticoagulants for the preservation of blood, plasma and other blood products *in vitro*. The compounds according to the invention are furthermore used for illnesses in which blood coagulation makes a crucial contribution to the course of the illness or represents a source of secondary pathology, such as, for example, in cancer, including metastasis, inflammatory disorders, including arthritis, and diabetes.
- In the treatment of the illnesses described, the compounds according to the invention are also employed in combination with other thrombolytically

active compounds, such as, for example, with "tissue plasminogen activator" t-PA, modified t-PA, streptokinase or urokinase. The compounds according to the invention are given either at the same time as or before or after the other substances mentioned.

5 Particular preference is given to simultaneous administration with aspirin in order to prevent recurrence of the clot formation.

The compounds according to the invention are also used in combination with blood platelet glycoprotein receptor (IIb/IIIa) antagonists, which inhibit
10 blood platelet aggregation.

The invention relates to the compounds of the formula I and their salts and to a process for the preparation of compounds of the formula I according to
15 Claim 1 and their salts, characterised in that

- a) they are liberated from one of their functional derivatives by treatment with a solvolysing or hydrogenolysing agent by
 - i) liberating an amidino group from their hydroxyl, oxadiazole or oxazolidinone derivative by hydrogenolysis or solvolysis,
 - ii) replacing a conventional amino-protecting group by hydrogen by treatment with a solvolysing or hydrogenolysing agent, or liberating an amino group protected by a conventional protecting group,
- 30 or
 - b) converting a cyano group into an amidino group ,
- 35 or

c) reacting a compound of the formula II

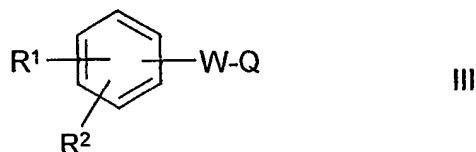
5 Z-Y-T II

in which

Z is HNR^3- or $\text{HNR}^3\text{C}(\text{R}^4)_2-$

10 and R^3 , R^4 , Y and T are as defined in Claim 1,

15 with a compound of the formula III

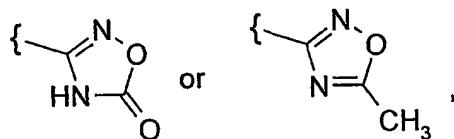


in which

20

R^1 is $-\text{C}(\text{=NH})-\text{NH}_2$ which is monosubstituted by $\text{C}(\text{=O})\text{R}^3$, COOR^3 ,
OR³ or by a conventional amino-protecting group, or is

25



30

Q is $-\text{CO-L}$ or $-\text{C}(\text{R}^4)_2\text{-L}$,
L is Cl, Br, I or a free or reactively functionally modified OH group,
and

35 and R^2 , R^3 , R^4 and W are as defined in Claim 1,

and/or

d) converting a base or acid of the formula I into one of its salts.

5 The invention also relates to the optically active forms (stereoisomers), the enantiomers, the racemates, the diastereomers and the hydrates and solvates of these compounds. The term solvates of the compounds is taken to mean adductions of inert solvent molecules onto the compounds which form owing to their mutual attractive force. Solvates are, for
10 example, monohydrates or dihydrates or alcoholates.

The term pharmaceutically usable derivatives is taken to mean, for example, the salts of the compounds according to the invention and also
15 so-called prodrug compounds.

The term prodrug derivatives is taken to mean compounds of the formula I which have been modified with, for example, alkyl or acyl groups, sugars or oligopeptides and which are rapidly cleaved in the organism to give the
20 effective compounds according to the invention.

These also include biodegradable polymer derivatives of the compounds according to the invention, as described, for example, in Int. J. Pharm.
115, 61-67 (1995).

25 The invention also relates to mixtures of the compounds of the formula I according to the invention, for example mixtures of two diastereomers, for example in the ratio 1:1, 1:2, 1:3, 1:4, 1:5, 1:10, 1:100 or 1:1000.

30 These are particularly preferably mixtures of stereoisomeric compounds.

35 The invention also relates, in particular, to the -C(=NH)-NH₂- compounds of the formula I which are substituted by -COA, -COOA, -OH or by a conventional amino-protecting group.

For all radicals which occur more than once, such as, for example, A, their meanings are independent of one another.

5 Above and below, the radicals or parameters W, X, Y, T, R¹ and R² are as defined under the formula I, unless expressly stated otherwise.

A is alkyl, is unbranched (linear) or branched, and has 1, 2, 3, 4, 5, 6, 7, 8,
9 or 10 carbon atoms. A is preferably methyl, furthermore ethyl, propyl, iso-
10 propyl, butyl, isobutyl, sec-butyl or tert-butyl, furthermore also pentyl, 1-, 2-
or 3-methylbutyl, 1,1-, 1,2- or 2,2-dimethylpropyl, 1-ethylpropyl, hexyl, 1-,
2-, 3- or 4-methylpentyl, 1,1-, 1,2-, 1,3-, 2,2-, 2,3- or 3,3-dimethylbutyl,
15 1- or 2-ethylbutyl, 1-ethyl-1-methylpropyl, 1-ethyl-2-methylpropyl, 1,1,2- or
1,2,2-trimethylpropyl, furthermore preferably, for example, trifluoromethyl.
A is very particularly preferably alkyl having 1-6 carbon atoms, preferably
methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl,
hexyl or trifluoromethyl.

20 Cycloalkyl is preferably cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl or
cycloheptyl.

Alkylene is preferably methylene, ethylene, propylene, butylene, pentylene
25 or hexylene, furthermore branched alkylene.

-COR³ (acyl) is preferably formyl, acetyl, propionyl, furthermore also
butyryl, pentanoyl, hexanoyl or, for example, benzoyl.

Ph is phenyl, Me is methyl, Et is ethyl, BOC is tert-butoxycarbonyl.

30 Hal is preferably F, Cl or Br, but alternatively I.

If R¹ is CON(R³)₂ or -[C(R⁴)_nN(R³)₂], CONH₂, NH₂ or CH₂NH₂ is preferred.

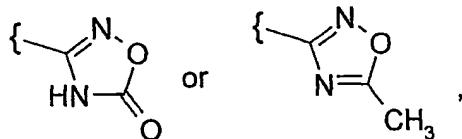
R¹ is particularly preferably CN, CONH₂, CONA₂, NH₂, CH₂NH₂,

CH₂CH₂NH₂,

35

-C(=NH)-NH₂ which is unsubstituted or monosubstituted by OH, OCOA or OCOOA, or is

5



where A is preferably alkyl having 1, 2, 3 or 4 carbon atoms.

10

R² is preferably H or F.

10

R³ is preferably H, A or -(CH₂)_n-Ar, particularly preferably, for example, H, alkyl having 1-6 carbon atoms, phenyl or benzyl.

15

W is preferably -C(R³)₂⁻, -OC(R³)₂⁻ or -NR³C(R³)₂⁻, particularly preferably, for example, -OCHR³ or -NHCHR³, where R³ is H, alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms, in which 3 H atoms may be replaced by F, or phenyl, benzyl or thienyl, each of which is unsubstituted or monosubstituted or disubstituted by F, methyl or ethyl.

20

X is preferably CONH or CONH(CH₂)₂⁻, particularly preferably CONH or CONHCH₂, furthermore CH₂O- or CH₂OCH₂.

25

Y is preferably alkylene or Ar-diyl, particularly preferably methylene, ethylene, propylene, or 1,4-phenylene which is unsubstituted or monosubstituted by F, ethoxycarbonylmethoxy or carboxymethoxy, furthermore alternatively pyridinedyl, preferably pyridine-2,5-diyl.

30

T is preferably N(R³)₂ or a monocyclic or bicyclic, saturated, unsaturated or aromatic heterocyclic radical having 1 to 2 N and/or O atoms, which is unsubstituted or monosubstituted or disubstituted by A and/or carbonyl oxygen, where R³ is H or A.

35

T is particularly preferably, for example, dimethylamino, diethylamino, morpholin-4-yl, 2-oxopiperidin-1-yl, 2-oxopyrrolidin-1-yl, 5,5-dimethyl-2-oxopyrrolidin-1-yl, 2-oxo-1*H*-pyridin-1-yl, 3-oxomorpholin-4-yl, 4-oxo-1*H*-pyridin-1-yl, 2,6-dioxopiperidin-1-yl, 2-oxopiperazin-1-yl, 2,5-dioxo-5-pyrrolidin-1-yl, 2-oxo-1,3-oxazolidin-3-yl, 2-azabicyclo[2.2.2]octan-3-on-2-yl or 3-oxo-2*H*-pyridazin-2-yl.

Ar is, for example, phenyl which is unsubstituted or monosubstituted, 10 disubstituted or trisubstituted by Hal, A, OR⁴, -[C(R⁴)₂]_n-COOR³ or O-[C(R⁴)₂]_o-COOR³.

Ar is preferably unsubstituted phenyl, naphthyl or biphenyl, furthermore 15 preferably phenyl, naphthyl or biphenyl, each of which is monosubstituted, disubstituted or trisubstituted, for example, by A, fluorine, chlorine, bromine, iodine, hydroxyl, methoxy, ethoxy, propoxy, butoxy, pentyloxy, hexyloxy, nitro, cyano, formyl, acetyl, propionyl, trifluoromethyl, amino, methylamino, ethylamino, dimethylamino, diethylamino, benzyloxy, sulfonamido, methylsulfonamido, ethylsulfonamido, propylsulfonamido, 20 butylsulfonamido, dimethylsulfonamido, phenylsulfonamido, carboxyl, methoxycarbonyl, ethoxycarbonyl, ethoxycarbonylmethoxy, carboxy- methoxy or aminocarbonyl.

Ar is particularly preferably, for example, phenyl which is unsubstituted or 25 monosubstituted or disubstituted by Hal, A, methoxy, methoxycarbonyl-methoxy, carboxymethoxy or ethoxycarbonylmethoxy.

Het is preferably, for example, 2- or 3-furyl, 2- or 3-thienyl, 1-, 2- or 30 3-pyrrolyl, 1-, 2-, 4- or 5-imidazolyl, 1-, 3-, 4- or 5-pyrazolyl, 2-, 4- or 5-oxazolyl, 3-, 4- or 5-isoxazolyl, 2-, 4- or 5-thiazolyl, 3-, 4- or 5-isothiazolyl, 2-, 3- or 4-pyridyl, 2-, 4-, 5- or 6-pyrimidinyl, furthermore preferably 35 1,2,3-triazol-1-, -4- or -5-yl, 1,2,4-triazol-1-, -3- or -5-yl, 1- or 5-tetrazolyl, 1,2,3-oxadiazol-4- or -5-yl, 1,2,4-oxadiazol-3- or -5-yl, 1,3,4-thiadiazol-2- or -5-yl, 1,2,4-thiadiazol-3- or -5-yl, 1,2,3-thiadiazol-4- or -5-yl, 3- or

4-pyridazinyl, pyrazinyl, 1-, 2-, 3-, 4-, 5-, 6- or 7-indolyl, 4- or 5-isoindolyl, 1-, 2-, 4- or 5-benzimidazolyl, 1-, 3-, 4-, 5-, 6- or 7-benzopyrazolyl, 2-, 4-, 5-, 6- or 7-benzoxazolyl, 3-, 4-, 5-, 6- or 7-benzisoxazolyl, 2-, 4-, 5-, 6- or 7-benzothiazolyl, 2-, 4-, 5-, 6- or 7-benzisothiazolyl, 4-, 5-, 6- or 7-benz-
5 2,1,3-oxadiazolyl, 2-, 3-, 4-, 5-, 6-, 7- or 8-quinolyl, 1-, 3-, 4-, 5-, 6-, 7- or 8-isoquinolyl, 3-, 4-, 5-, 6-, 7- or 8-quinolinyl, 2-, 4-, 5-, 6-, 7- or 8-quinazo-
10 linyl, 5- or 6-quinoxaliny, 2-, 3-, 5-, 6-, 7- or 8-2H-benzo-1,4-oxazinyl, furthermore preferably 1,3-benzodioxol-5-yl, 1,4-benzodioxan-6-yl, 2,1,3-
benzothiadiazol-4- or -5-yl or 2,1,3-benzoxadiazol-5-yl.

The heterocyclic radicals may also be partially or fully hydrogenated. Het can thus, for example, also be 2,3-dihydro-2-, -3-, -4- or -5-furyl, 2,5-dihydro-2-, -3-, -4- or -5-furyl, tetrahydro-2- or -3-furyl, 1,3-dioxolan-4-yl, 15 tetrahydro-2- or -3-thienyl, 2,3-dihydro-1-, -2-, -3-, -4- or -5-pyrrolyl, 2,5-dihydro-1-, -2-, -3-, -4- or -5-pyrrolyl, 1-, 2- or 3-pyrrolidinyl, tetrahydro-1-, -2- or -4-imidazolyl, 2,3-dihydro-1-, -2-, -3-, -4- or -5-pyrazolyl, tetrahydro-1-, -3- or -4-pyrazolyl, 1,4-dihydro-1-, -2-, -3- or -4-pyridyl, 1,2,3,4-tetrahydro-
20 1-, -2-, -3-, -4-, -5- or -6-pyridyl, 1-, 2-, 3- or 4-piperidinyl, 2-, 3- or 4-morpholinyl, tetrahydro-2-, -3- or -4-pyranyl, 1,4-dioxanyl, 1,3-dioxan-2-, -4- or -5-yl, hexahydro-1-, -3- or -4-pyridazinyl, hexahydro-1-, -2-, -4- or -5-pyrimidinyl, 1-, 2- or 3-piperazinyl, 1,2,3,4-tetrahydro-1-, -2-, -3-, -4-, -5-,
25 -6-, -7- or -8-quinolyl, 1,2,3,4-tetrahydro-1-, -2-, -3-, -4-, -5-, -6-, -7- or -8-isoquinolyl, 2-, 3-, 5-, 6-, 7- or 8-3,4-dihydro-2H-benzo-1,4-oxazinyl, furthermore preferably 2,3-methylenedioxyphenyl, 3,4-methylenedioxy-
30 phenyl, 2,3-ethylenedioxyphenyl, 3,4-ethylenedioxyphenyl, 3,4-(difluoro-methylenedioxy)phenyl, 2,3-dihydrobenzofuran-5- or -6-yl, 2,3-(2-oxo-methylenedioxy)phenyl or alternatively 3,4-dihydro-2H-1,5-benzodioxepin-6- or -7-yl, furthermore preferably 2,3-dihydrobenzofuranyl or 2,3-dihydro-2-oxofuranyl.

Het is very particularly preferably a monocyclic or bicyclic, saturated, unsaturated or aromatic heterocyclic radical having 1 to 2 N or O atoms which is unsubstituted or monosubstituted or disubstituted by carbonyl oxygen, such as, for example, morpholin-4-yl, 2-oxopiperidin-1-yl, 2-oxo-pyrrolidin-1-yl, 5,5-dimethyl-2-oxopyrrolidin-1-yl, 2-oxo-1*H*-pyridin-1-yl, 3-oxomorpholin-4-yl, 4-oxo-1*H*-pyridin-1-yl, 2,6-dioxopiperidin-1-yl, 2-oxo-piperazin-1-yl, 2,5-dioxopyrrolidin-1-yl, 2-oxo-1,3-oxazolidin-3-yl, 3-oxo-2*H*-pyridazin-2-yl, 2-caprolactam-1-yl, 2-oxo-1,3-oxazinan-3-yl or 2-aza-
5 bicyclo[2.2.2]octan-3-on-2-yl.

Het' is preferably, for example, 2- or 3-furyl, 2- or 3-thienyl, 1-, 2- or 3-pyrrolyl, 1-, 2-, 4- or 5-imidazolyl, 1-, 3-, 4- or 5-pyrazolyl, 2-, 4- or 5-oxazolyl, 3-, 4- or 5-isoxazolyl, 2-, 4- or 5-thiazolyl, 3-, 4- or 5-isothiazolyl, 2-, 3- or 4-pyridyl, 2-, 4-, 5- or 6-pyrimidinyl, furthermore preferably
10 1,2,3-triazol-1-, -4- or -5-yl, 1,2,4-triazol-1-, -3- or -5-yl, 1- or 5-tetrazolyl, 1,2,3-oxadiazol-4- or -5-yl, 1,2,4-oxadiazol-3- or -5-yl, 1,3,4-thiadiazol-2- or -5-yl, 1,2,4-thiadiazol-3- or -5-yl, 1,2,3-thiadiazol-4- or -5-yl, 3- or
15 4-pyridazinyl, pyrazinyl, 1-, 2-, 3-, 4-, 5-, 6- or 7-indolyl, 4- or 5-isoindolyl, 1-, 2-, 4- or 5-benzimidazolyl, 1-, 3-, 4-, 5-, 6- or 7-benzopyrazolyl, 2-, 4-, 5-, 6- or 7-benzoxazolyl, 3-, 4-, 5-, 6- or 7-benzisoxazolyl, 2-, 4-, 5-, 6- or 7-benzothiazolyl, 2-, 4-, 5-, 6- or 7-benzisothiazolyl, 4-, 5-, 6- or 7-benz-
20 2,1,3-oxadiazolyl, 2-, 3-, 4-, 5-, 6-, 7- or 8-quinolyl, 1-, 3-, 4-, 5-, 6-, 7- or 8-isoquinolyl, 3-, 4-, 5-, 6-, 7- or 8-quinolinyl, 2-, 4-, 5-, 6-, 7- or 8-quinazolinyl, 5- or 6-quinoxalinyl, 2-, 3-, 5-, 6-, 7- or 8-2*H*-benzo-1,4-oxazinyl, furthermore preferably 1,3-benzodioxol-5-yl, 1,4-benzodioxan-6-yl, 2,1,3-
25 benzothiadiazol-4- or -5-yl or 2,1,3-benzoxadiazol-5-yl.

The heterocyclic radicals may also be partially or fully hydrogenated.
30 Het' can thus, for example, also be 2,3-dihydro-2-, -3-, -4- or -5-furyl, 2,5-dihydro-2-, -3-, -4- or -5-furyl, tetrahydro-2- or -3-furyl, 1,3-dioxolan-4-yl, tetrahydro-2- or -3-thienyl, 2,3-dihydro-1-, -2-, -3-, -4- or -5-pyrrolyl, 2,5-dihydro-1-, -2-, -3-, -4- or -5-pyrrolyl, 1-, 2- or 3-pyrrolidinyl, tetrahydro-1-, -35

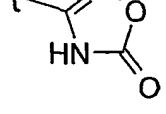
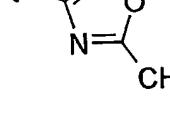
2- or -4-imidazolyl, 2,3-dihydro-1-, -2-, -3-, -4- or -5-pyrazolyl, tetrahydro-1-,
-3- or -4-pyrazolyl, 1,4-dihydro-1-, -2-, -3- or -4-pyridyl, 1,2,3,4-tetrahydro-
1-, -2-, -3-, -4-, -5- or -6-pyridyl, 1-, 2-, 3- or 4-piperidinyl, 2-, 3- or 4-
morpholinyl, tetrahydro-2-, -3- or -4-pyranyl, 1,4-dioxanyl, 1,3-dioxan-2-,
5 -4- or -5-yl, hexahydro-1-, -3- or -4-pyridazinyl, hexahydro-1-, -2-, -4- or -5-
pyrimidinyl, 1-, 2- or 3-piperazinyl, 1,2,3,4-tetrahydro-1-, -2-, -3-, -4-, -5-,
-6-, -7- or -8-quinolyl, 1,2,3,4-tetrahydro-1-, -2-, -3-, -4-, -5-, -6-, -7- or -8-
isoquinolyl, 2-, 3-, 5-, 6-, 7- or 8-3,4-dihydro-2H-benzo-1,4-oxazinyl,
10 furthermore preferably 2,3-methylenedioxyphenyl, 3,4-methylenedioxy-
phenyl, 2,3-ethylenedioxyphenyl, 3,4-ethylenedioxyphenyl, 3,4-(difluoro-
methylenedioxy)phenyl, 2,3-dihydrobenzofuran-5- or -6-yl, 2,3-(2-oxo-
methylenedioxy)phenyl or alternatively 3,4-dihydro-2H-1,5-benzodioxepin-
15 6- or -7-yl, furthermore preferably 2,3-dihydrobenzofuranyl or 2,3-dihydro-
2-oxofuranyl.

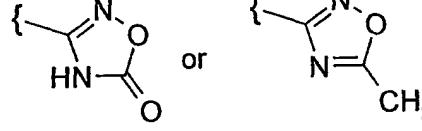
m is preferably 2, furthermore alternatively 0 or 1.
20 n is preferably 1, furthermore alternatively 0 or 2.

The compounds of the formula I may have one or more chiral centres and
therefore occur in various stereoisomeric forms. The formula I covers all
25 these forms.

Accordingly, the invention relates in particular to the compounds of the
formula I in which at least one of the said radicals has one of the preferred
30 meanings indicated above. Some preferred groups of compounds may be
expressed by the following sub-formulae Ia to Im, which conform to the
formula I and in which the radicals not designated in greater detail are as
defined under the formula I, but in which

35 in Ia R² is H;

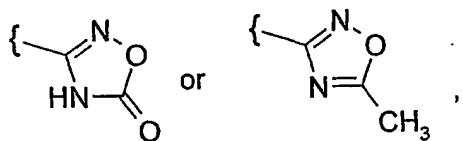
in Ib	<p>R^1 is $-C(=NH)-NH_2$ which is unsubstituted or monosubstituted by OH, or is</p> <p style="text-align: center;">5 {  or {  ;</p>
10 in Ic	<p>Ar is phenyl which is unsubstituted or monosubstituted or disubstituted by Hal, A, OR⁴ or O-[C(R⁴)₂]-COOR³;</p>
15 in Id	<p>Het is a monocyclic or bicyclic, saturated, unsaturated or aromatic heterocyclic radical having 1 to 2 N and/or O atoms which is monosubstituted or disubstituted by A and/or carbonyl oxygen;</p>
20 in Ie	<p>W is $-OC(R^3)_2-$ or $-NR^3C(R^3)_2-$;</p>
25 in If	<p>W is $-OC(R^3)_2-$ or $-NR^3C(R^3)_2-$, R^3 is H, A or $-(CH_2)_n-Ar$,</p> <p>Ar is phenyl which is unsubstituted or monosubstituted or disubstituted by Hal, A, OR⁴ or O-[C(R⁴)₂]-COOR³,</p> <p>n is 0 or 1;</p>
30 in Ig	<p>W is $-OCHR^3-$ or $-NHCHR^3-$, R^3 is H, A or $-(CH_2)_n-Ar$,</p> <p>Ar is phenyl which is unsubstituted or monosubstituted or disubstituted by Hal, A, OR⁴ or O-[C(R⁴)₂]-COOR³,</p> <p>n is 0 or 1;</p>
35 in Ih	<p>X is CONH or CONH(CH₂)₂-;</p>

5	in II	Y is alkylene or Ar-diyl, Ar is phenyl which is unsubstituted or monosubstituted or disubstituted by Hal, A, OR ⁴ or O-[C(R ⁴) ₂] _o -COOR ³ ;
10	in Ij	T is N(R ³) ₂ or a monocyclic or bicyclic, saturated, unsaturated or aromatic heterocyclic radical having 1 to 2 N and/or O atoms which is monosubstituted or disubstituted by A and/or carbonyl oxygen, R ³ is H or A;
15	in Ik	R ¹ is -C(=NH)-NH ₂ which is unsubstituted or monosubstituted by OH, or is 
20		R ² is H, R ³ is H, A or -(CH ₂) _n -Ar, R ⁴ is H or A, W is -OC(R ³) ₂ - or -NR ³ C(R ³) ₂ -,
25		X is CONH or CONH(CH ₂) ₂ , Y is alkylene or Ar-diyl, T is N(R ³) ₂ or a monocyclic or bicyclic, saturated, unsaturated or aromatic heterocyclic radical having 1 to 2 N and/or O atoms which is monosubstituted or disubstituted by A and/or carbonyl oxygen,
30		R ³ is H or A, Ar is phenyl which is unsubstituted or monosubstituted or disubstituted by Hal, A, OR ⁴ or O-[C(R ⁴) ₂] _o -COOR ³ ,
35		

A is unbranched or branched alkyl having 1-6 carbon atoms,
 in which 1-7 H atoms may be replaced by F,
 n is 0 or 1;

5 in II R¹ is -C(=NH)-NH₂ which is unsubstituted or monosubstituted
 by OH, or is

10



15

R² is H,
 R³ is H, A or -(CH₂)_n-Ar,
 R⁴ is H or A,
 W is -OC(R³)₂- or -NR³C(R³)₂-,
 X is CONH or CONH(CH₂)₂,

20

Y is alkylene or Ar-diyl,
 T is dimethylamino, diethylamino, morpholin-4-yl, 2-oxo-piperidin-1-yl, 2-oxopyrrolidin-1-yl, 5,5-dimethyl-2-oxo-pyrrolidin-1-yl, 2-oxo-1*H*-pyridin-1-yl, 3-oxomorpholin-4-yl, 4-oxo-1*H*-pyridin-1-yl, 2,6-dioxopiperidin-1-yl, 2-oxo-piperazin-1-yl, 2,5-dioxopyrrolidin-1-yl, 2-oxo-1,3-oxazolidin-3-yl, 2-azabicyclo[2.2.2]octan-3-on-2-yl or 3-oxo-2*H*-pyridazin-2-yl,

25

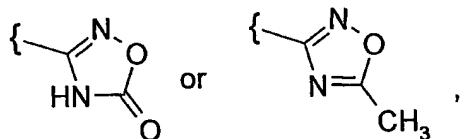
Ar is phenyl which is unsubstituted or monosubstituted or disubstituted by Hal, A, OR⁴ or O-[C(R⁴)₂]-COOR³,

30

A is unbranched or branched alkyl having 1-6 carbon atoms,
 in which 1-7 H atoms may be replaced by F,
 n is 0 or 1;

in Im R¹ is CN, NH₂, CONA₂, CH₂NH₂, CH₂CH₂NH₂, -C(=NH)-NH₂
 which is unsubstituted or monosubstituted by OH, COOR³,
 OCOA or OCOOA, or is

5



10

R² is H or F,R³ is H, A or -(CH₂)_n-Ar or thiienyl,R⁴ is H or A,W is -C(R³)₂⁻, -OC(R³)₂⁻ or -NR³C(R³)₂⁻,X is CONH, CONH(CH₂), CONH(CH₂)₂, CH₂O- or CH₂OCH₂,

Y is alkylene, Ar-diyl or pyridinediyl,

15

T is dimethylamino, diethylamino, morpholin-4-yl,

2-oxopiperidin-1-yl, 2-oxopyrrolidin-1-yl, 5,5-dimethyl-2-oxopyrrolidin-1-yl, 2-oxo-1*H*-pyridin-1-yl, 3-oxomorpholin-4-yl, 4-oxo-1*H*-pyridin-1-yl, 2,6-dioxopiperidin-1-yl, 2-oxo-piperazin-1-yl, 2,5-dioxopyrrolidin-1-yl, 2-oxo-1,3-oxazolidin-3-yl, 3-oxo-2*H*-pyridazin-2-yl, 2-caprolactam-1-yl, 2-oxo-1,3-oxazinan-3-yl or 2-azabicyclo[2.2.2]octan-3-on-2-yl,

20

25

Ar is phenyl which is unsubstituted or monosubstituted or

disubstituted by Hal, CF₃, A, OA, methoxycarbonyl-methoxy, ethoxycarbonylmethoxy or carboxymethoxy,

A is unbranched or branched alkyl having 1-6 carbon atoms,

30

in which 1-7 H atoms may be replaced by F,

n is 0 or 1;

and their pharmaceutically usable derivatives, solvates and stereoisomers,
 35 including mixtures thereof in all ratios.

The compounds of the formula I and also the starting materials for their preparation are, in addition, prepared by methods known per se, as described in the literature (for example in the standard works, such as Houben-Weyl, Methoden der organischen Chemie [Methods of Organic Chemistry], Georg-Thieme-Verlag, Stuttgart), to be precise under reaction conditions which are known and suitable for the said reactions. Use can also be made here of variants which are known per se, but are not mentioned here in greater detail.

10

If desired, the starting materials can also be formed in situ so that they are not isolated from the reaction mixture, but instead are immediately converted further into the compounds of the formula I.

15

Compounds of the formula I can preferably be obtained by liberating compounds of the formula I from one of their functional derivatives by treatment with a solvolysing or hydrogenolysing agent.

20

Preferred starting materials for the solvolysis or hydrogenolysis are those which conform to the formula I, but contain corresponding protected amino and/or hydroxyl groups instead of one or more free amino and/or hydroxyl groups, preferably those which carry an amino-protecting group instead of an H atom bonded to an N atom, in particular those which carry an R'-N group, in which R' is an amino-protecting group, instead of an HN group, and/or those which carry a hydroxyl-protecting group instead of the H atom of a hydroxyl group, for example those which conform to the formula I, but carry a -COOR" group, in which R" is a hydroxyl-protecting group, instead of a -COOH group.

25

Preferred starting materials are also the oxadiazole derivatives, which can be converted into the corresponding amidino compounds.

30

35

The amidino group can be liberated from its oxadiazole derivative by, for example, treatment with hydrogen in the presence of a catalyst (for example Raney nickel). Suitable solvents are those indicated below, in particular alcohols, such as methanol or ethanol, organic acids, such as acetic acid or propionic acid, or mixtures thereof. The hydrogenolysis is generally carried out at temperatures between about 0 and 100° and pressures between about 1 and 200 bar, preferably at 20-30° (room temperature) and 1-10 bar.

10

The oxadiazole group is introduced, for example, by reaction of the cyano compounds with hydroxylamine and reaction with phosgene, dialkyl carbonate, chloroformic acid esters, N,N'-carbonyldiimidazole or acetic anhydride.

15

It is also possible for a plurality of – identical or different – protected amino and/or hydroxyl groups to be present in the molecule of the starting material. If the protecting groups present are different from one another, they can in many cases be cleaved off selectively.

20

The term "amino-protecting group" is known in general terms and relates to groups which are suitable for protecting (blocking) an amino group against chemical reactions, but which are easy to remove after the desired chemical reaction has been carried out elsewhere in the molecule. Typical of such groups are, in particular, unsubstituted or substituted acyl, aryl, aralkoxymethyl or aralkyl groups. Since the amino-protecting groups are removed after the desired reaction (or reaction sequence), their type and size are furthermore not crucial; however, preference is given to those having 1-20, in particular 1-8, carbon atoms. The term "acyl group" is to be understood in the broadest sense in connection with the present process.

30

35 It includes acyl groups derived from aliphatic, araliphatic, aromatic or

heterocyclic carboxylic acids or sulfonic acids, and, in particular, alkoxy-carbonyl, aryloxycarbonyl and especially aralkoxycarbonyl groups.

Examples of such acyl groups are alkanoyl, such as acetyl, propionyl and butyryl; aralkanoyl, such as phenylacetyl; aroyl, such as benzoyl and tolyl; aryloxalkanoyl, such as POA; alkoxy carbonyl, such as methoxycarbonyl, ethoxycarbonyl, 2,2,2-trichloroethoxycarbonyl, BOC (tert-butoxycarbonyl) and 2-iodoethoxycarbonyl; aralkoxycarbonyl, such as CBZ ("carbo-benzoxy"), 4-methoxybenzyloxycarbonyl and FMOC; and arylsulfonyl, such as Mtr. Preferred amino-protecting groups are BOC and Mtr, furthermore CBZ, Fmoc, benzyl and acetyl.

The term "hydroxyl-protecting group" is likewise known in general terms and relates to groups which are suitable for protecting a hydroxyl group against chemical reactions, but are easily removable after the desired chemical reaction has been carried out elsewhere in the molecule. Typical of such groups are the above-mentioned unsubstituted or substituted aryl, aralkyl or acyl groups, furthermore also alkyl groups. The nature and size of the hydroxyl-protecting groups are not crucial since they are removed again after the desired chemical reaction or reaction sequence; preference is given to groups having 1-20, in particular 1-10, carbon atoms. Examples of hydroxyl-protecting groups are, inter alia, benzyl, 4-methoxybenzyl, p-nitrobenzoyl, p-toluenesulfonyl, tert-butyl and acetyl, where benzyl and tert-butyl are particularly preferred.

The compounds of the formula I are liberated from their functional derivatives – depending on the protecting group used – for example using strong acids, advantageously using TFA or perchloric acid, but also using other strong inorganic acids, such as hydrochloric acid or sulfuric acid, strong organic carboxylic acids, such as trichloroacetic acid, or sulfonic acids, such as benzene- or p-toluenesulfonic acid. The presence of an

additional inert solvent is possible, but is not always necessary. Suitable inert solvents are preferably organic, for example carboxylic acids, such as acetic acid, ethers, such as tetrahydrofuran or dioxane, amides, such as DMF, halogenated hydrocarbons, such as dichloromethane, furthermore 5 also alcohols, such as methanol, ethanol or isopropanol, and water.

Mixtures of the above-mentioned solvents are furthermore suitable. TFA is 10 preferably used in excess without addition of a further solvent, and perchloric acid is preferably used in the form of a mixture of acetic acid and 70% perchloric acid in the ratio 9:1. The reaction temperatures for the cleavage are advantageously between about 0 and about 50°, preferably between 15 and 30° (room temperature).

15 The BOC, O-but and Mtr groups can, for example, preferably be cleaved off using TFA in dichloromethane or using approximately 3 to 5N HCl in dioxane at 15-30°, and the Fmoc group can be cleaved off using an approximately 5 to 50% solution of dimethylamine, diethylamine or 20 piperidine in DMF at 15-30°.

Protecting groups which can be removed hydrogenolytically (for example 25 CBZ, benzyl or the liberation of the amidino group from its oxadiazole derivative) can be cleaved off, for example, by treatment with hydrogen in the presence of a catalyst (for example a noble-metal catalyst, such as palladium, advantageously on a support, such as carbon). Suitable solvents here are those indicated above, in particular, for example, 30 alcohols, such as methanol or ethanol, or amides, such as DMF. The hydrogenolysis is generally carried out at temperatures between about 0 and 100° and pressures between about 1 and 200 bar, preferably at 20-30° and 1-10 bar. Hydrogenolysis of the CBZ group succeeds well, for 35 example, on 5 to 10% Pd/C in methanol or using ammonium formate (instead of hydrogen) on Pd/C in methanol/DMF at 20-30°.

- Examples of suitable inert solvents are hydrocarbons, such as hexane, petroleum ether, benzene, toluene or xylene; chlorinated hydrocarbons, such as trichloroethylene, 1,2-dichloroethane, tetrachloromethane,
5 trifluoromethylbenzene, chloroform or dichloromethane; alcohols, such as methanol, ethanol, isopropanol, n-propanol, n-butanol or tert-butanol; ethers, such as diethyl ether, diisopropyl ether, tetrahydrofuran (THF) or dioxane; glycol ethers, such as ethylene glycol monomethyl or monoethyl
10 ether or ethylene glycol dimethyl ether (diglyme); ketones, such as acetone or butanone; amides, such as acetamide, dimethylacetamide, N-methylpyrrolidone (NMP) or dimethylformamide (DMF); nitriles, such as acetonitrile; sulfoxides, such as dimethyl sulfoxide (DMSO); carbon
15 disulfide; carboxylic acids, such as formic acid or acetic acid; nitro compounds, such as nitromethane or nitrobenzene; esters, such as ethyl acetate, or mixtures of the said solvents.
- 20 A cyano group is converted into an amidino group by reaction with, for example, hydroxylamine followed by reduction of the N-hydroxyamidine using hydrogen in the presence of a catalyst, such as, for example, Pd/C. In order to prepare an amidine of the formula I, it is also possible to adduct
25 ammonia onto a nitrile. The addition is preferably carried out in a number of steps by, in a manner known per se, a) converting the nitrile into a thioamide using H₂S, converting the thioamide into the corresponding S-alkylimidothioester using an alkylating agent, for example CH₃I, and reacting the thioester in turn with NH₃ to give the amidine, b) converting the
30 nitrile into the corresponding imidoester using an alcohol, for example ethanol in the presence of HCl, and treating the imidoester with ammonia (Pinner synthesis), or c) reacting the nitrile with lithium bis(trimethylsilyl)amide, and subsequently hydrolysing the product.
35

Esters can be saponified, for example, using acetic acid or using NaOH or KOH in water, water/THF or water/dioxane, at temperatures between 0 and 100°.

5 Free amino groups can furthermore be acylated in a conventional manner using an acid chloride or anhydride or alkylated using an unsubstituted or substituted alkyl halide, or reacted with CH₃-C(=NH)-OEt, advantageously in an inert solvent, such as dichloromethane or THF and/or in the presence
10 of a base, such as triethylamine or pyridine, at temperatures between -60 and +30°.

If desired, the starting materials can also be formed in situ so that they are
15 not isolated from the reaction mixture, but instead are immediately converted further into the compounds of the formula I.

Compounds of the formula I in which R¹ is in protected form can preferably
20 be obtained by reacting compounds of the formula II with compounds of the formula III.

The reaction is generally carried out in an inert solvent, in the presence of an acid-binding agent, preferably an alkali or alkaline earth metal
25 hydroxide, carbonate or bicarbonate, or in the presence of another salt of a weak acid of the alkali or alkaline earth metals, preferably of potassium, sodium, calcium or caesium. The addition of an organic base, such as triethylamine, dimethylaniline, pyridine or quinoline, may also be favourable. Depending on the conditions used, the reaction time is
30 between a few minutes and 14 days, and the reaction temperature is between about 0° and 150°, normally between 20° and 130°.

35 Examples of suitable inert solvents are water; hydrocarbons, such as hexane, petroleum ether, benzene, toluene or xylene; chlorinated

hydrocarbons, such as trichloroethylene, 1,2-dichloroethane, carbon tetrachloride, chloroform or dichloromethane; alcohols, such as methanol, ethanol, isopropanol, n-propanol, n-butanol or tert-butanol; ethers, such as diethyl ether, diisopropyl ether, tetrahydrofuran (THF) or dioxane; glycol ethers, such as ethylene glycol monomethyl or monoethyl ether or ethylene glycol dimethyl ether (diglyme); ketones, such as acetone or butanone; amides, such as acetamide, dimethylacetamide or dimethylformamide (DMF); nitriles, such as acetonitrile; sulfoxides, such as dimethyl sulfoxide (DMSO); carbon disulfide; carboxylic acids, such as formic acid or acetic acid; nitro compounds, such as nitromethane or nitrobenzene; esters, such as ethyl acetate, or mixtures of the said solvents.

15

The starting compounds of the formulae II and III are generally known. If they are novel, however, they can be prepared by methods known per se.

20

In the compounds of the formula III, L is preferably Cl, Br, I or a reactively modified OH group, such as, for example, an activated ester, an imidazolide or alkylsulfonyloxy having 1-6 carbon atoms (preferably methylsulfonyloxy or trifluoromethylsulfonyloxy) or arylsulfonyloxy having 6-10 carbon atoms (preferably phenyl- or p-tolylsulfonyloxy).

25

A base of the formula I can be converted into the associated acid-addition salt using an acid, for example by reaction of equivalent amounts of the base and the acid in an inert solvent, such as ethanol, followed by evaporation. Suitable acids for this reaction are, in particular, those which give physiologically acceptable salts. Thus, it is possible to use inorganic acids, for example sulfuric acid, nitric acid, hydrohalic acids, such as hydrochloric acid or hydrobromic acid, phosphoric acids, such as orthophosphoric acid, or sulfamic acid, furthermore organic acids, in particular

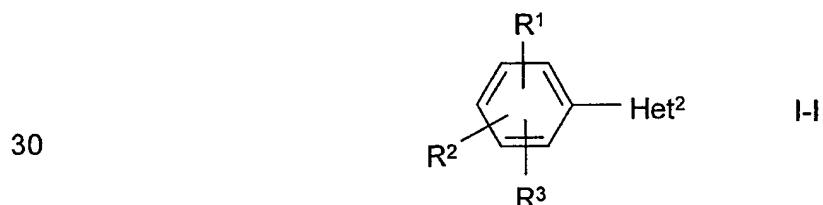
30

35

aliphatic, alicyclic, araliphatic, aromatic or heterocyclic monobasic or polybasic carboxylic, sulfonic or sulfuric acids, for example formic acid, acetic acid, propionic acid, pivalic acid, diethylacetic acid, malonic acid, succinic acid, pimelic acid, fumaric acid, maleic acid, lactic acid, tartaric acid, malic acid, citric acid, gluconic acid, ascorbic acid, nicotinic acid, 5 isonicotinic acid, methane- or ethanesulfonic acid, ethanedisulfonic acid, 2-hydroxyethanesulfonic acid, benzenesulfonic acid, p-toluenesulfonic acid, naphthalenemono- and -disulfonic acids, and laurylsulfuric acid. Salts with 10 physiologically unacceptable acids, for example picrates, can be used for the isolation and/or purification of the compounds of the formula I.

On the other hand, compounds of the formula I can be converted into the 15 corresponding metal salts, in particular alkali metal or alkaline earth metal salts, or into the corresponding ammonium salts using bases (for example sodium hydroxide, potassium hydroxide, sodium carbonate or potassium carbonate).
20 It is also possible to use physiologically acceptable organic bases, such as, for example, ethanolamine.

Some of the starting compounds are novel. The invention therefore 25 furthermore relates to the intermediates of the formula I-I



in which

R¹ is NO₂, CN, NHA, NHCOA, NACF₃, NH₂ or (CH₂)_nNH₂,
35 R² and
R³ are each, independently of one another, H, Hal, A, OA or CF₃,

- A is alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,
Het² is a monocyclic or bicyclic, saturated, unsaturated or aromatic
heterocyclic radical having from 1 to 4 N, O and/or S atoms which
is unsubstituted or monosubstituted, disubstituted or trisubstituted
5 by Hal, A, OA, CN, COOA, CONH₂, NHCOA, NSO₂A, S(O)_mA
and/or carbonyl oxygen,
n is 0, 1, 2 or 3,
and salts thereof.
10
In the compounds of the formula I-I, A is preferably methyl, ethyl, propyl,
isopropyl, butyl or tert-butyl.
Het² is preferably piperidinyl, pyrrolidinyl, piperazinyl, azepamyl, pyrazinyl,
15 3,4,5,6-tetrahydropyridinyl, pyridinyl, 2-azabicyclo[2.2.2]octanyl, 1,3-
oxazinanyl or oxazolidinyl.
The invention furthermore relates to the intermediates selected from the
20 group consisting of
1-(3-fluoro-4-nitrophenyl)piperidin-2-one,
1-(4-amino-3-fluorophenyl)piperidin-2-one,
25 *tert*-butyl 4-(4-nitrophenyl)-3-oxopiperazin-1-carboxylate,
tert-butyl 4-(4-aminophenyl)-3-oxopiperazin-1-carboxylate,
1-(4-nitrophenyl)azepam-2-one,
1-(4-aminophenyl)azepam-2-one,
30 1-(3-fluoro-4-nitrophenyl)azepam-2-one,
1-(4-amino-3-fluorophenyl)azepam-2-one,
1-(2-methyl-4-nitrophenyl)azepam-2-one,
1-(4-amino-2-methylphenyl)azepam-2-one,
35 1-(4-nitrophenyl)-1*H*-pyrazin-2-one,
1-(4-aminophenyl)-1*H*-pyrazin-2-one,

- 1-(2,5-dimethyl-4-nitrophenyl)piperidin-2-one,
1-(4-amino-2,5-dimethylphenyl)piperidin-2-one,
2,2,2-trifluoro-N-methyl-N-[4-(2-oxopiperidin-1-yl)phenyl]acetamide,
1-(4-methylaminophenyl)piperidin-2-one,
5 1-(3-methyl-4-nitrophenyl)piperidin-2-one,
1-(4-amino-3-methylphenyl)piperidin-2-one,
5'-nitro-3,4,5,6-tetrahydro-1,2'-bipyridinyl-2-one,
5'-amino-3,4,5,6-tetrahydro-1,2'-bipyridinyl-2-one,
10 2-(4-nitrophenyl)-2-azabicyclo[2.2.2]octan-3-one,
2-(4-aminophenyl)-2-azabicyclo[2.2.2]octan-3-one,
2-(4-nitro-2-trifluoromethylphenyl)-2-azabicyclo[2.2.2]octan-3-one,
2-(4-amino-2-trifluoromethylphenyl)-2-azabicyclo[2.2.2]octan-3-one,
15 2-(2-methyl-4-nitrophenyl)-2-azabicyclo[2.2.2]octan-3-one,
2-(4-amino-2-methylphenyl)-2-azabicyclo[2.2.2]octan-3-one,
2-(2-methoxy-4-nitrophenyl)-2-azabicyclo[2.2.2]octan-3-one,
2-(4-amino-2-methoxyphenyl)-2-azabicyclo[2.2.2]octan-3-one,
20 1-(2-methoxy-4-nitrophenyl)piperidin-2-one,
1-(4-amino-2-methoxyphenyl)piperidin-2-one,
1-(4-nitro-2-trifluoromethylphenyl)piperidin-2-one,
1-(4-amino-2-trifluoromethylphenyl)piperidin-2-one,
25 3-(2-methyl-4-nitrophenyl)-1,3-oxazinan-2-one,
3-(4-amino-2-methylphenyl)-1,3-oxazinan-2-one,
1-(2-chloro-4-nitrophenyl)pyrrolidin-2-one,
1-(4-amino-2-chlorophenyl)pyrrolidin-2-one,
30 1-(2-methoxy-4-nitrophenyl)pyrrolidine-2,5-dione,
1-(4-amino-2-methoxyphenyl)pyrrolidine-2,5-dione,
5,5-dimethyl-1-(4-nitrophenyl)pyrrolidin-2-one,
1-(4-aminophenyl)-5,5-dimethylpyrrolidin-2-one,
35 3-(2-methyl-4-nitrophenyl)oxazolidin-2-one,
3-(4-amino-2-methylphenyl)oxazolidin-2-one,

1-(2-fluoro-4-nitrophenyl)azepam-2-one,
1-(4-amino-2-fluorophenyl)azepam-2-one,
4-(2-oxo-2H-pyridin-1-yl)benzonitrile,
5 1-(4-aminomethylphenyl)-1*H*-pyridin-2-one,
1-(4-aminomethylphenyl)piperidin-2-one,
1-(2-ethyl-5-nitrophenyl)pyrrolidin-2-one,
1-(5-amino-2-ethylphenyl)pyrrolidin-2-one,
10 1-(4-aminophenyl)piperidine-2,6-dione,
and salts thereof.

Compounds of the formula I according to the invention may be chiral owing
to their molecular structure and may accordingly occur in various enantio-
15 meric forms. They can therefore exist in racemic or in optically active form.

Since the pharmaceutical activity of the racemates or stereoisomers of the
compounds according to the invention may differ, it may be desirable to
20 use the enantiomers. In these cases, the end product or even the interme-
diates can be separated into enantiomeric compounds by chemical or
physical measures known to the person skilled in the art or even employed
as such in the synthesis.

25 In the case of racemic amines, diastereomers are formed from the mixture
by reaction with an optically active resolving agent. Examples of suitable
resolving agents are optically active acids, such as the R and S forms of
tartaric acid, diacetyl tartaric acid, dibenzoyltartaric acid, mandelic acid,
30 malic acid, lactic acid, suitable N-protected amino acids (for example N-
benzoylproline or N-benzenesulfonylproline), or the various optically active
camphorsulfonic acids. Also advantageous is chromatographic enantiomer
resolution with the aid of an optically active resolving agent (for example
35 dinitrobenzoylphenylglycine, cellulose triacetate or other derivatives of
carbohydrates or chirally derivatised methacrylate polymers immobilised

on silica gel). Suitable eluents for this purpose are aqueous or alcoholic solvent mixtures, such as, for example, hexane/isopropanol/ acetonitrile, for example in the ratio 82:15:3.

5 The invention furthermore relates to the use of the compounds of the formula I and/or their physiologically acceptable salts for the preparation of pharmaceutical preparations, in particular by non-chemical methods. They can be converted here into a suitable dosage form together with at least
10 one solid, liquid and/or semi-liquid excipient or assistant and, if desired, in combination with one or more further active ingredients.

15 The invention furthermore relates to medicaments comprising at least one compound of the formula I and/or its pharmaceutically usable derivatives, solvates and stereoisomers, including mixtures thereof in all ratios, and, if desired, excipients and/or assistants.

20 These preparations can be used as medicaments in human or veterinary medicine. Suitable excipients are organic or inorganic substances which are suitable for enteral (for example oral), parenteral or topical administration and do not react with the novel compounds, for example water, vegetable oils, benzyl alcohols, alkylene glycols, polyethylene glycols, glycerol
25 triacetate, gelatin, carbohydrates, such as lactose or starch, magnesium stearate, talc or vaseline. Suitable for oral administration are, in particular, tablets, pills, coated tablets, capsules, powders, granules, syrups, juices or drops, suitable for rectal administration are suppositories, suitable for
30 parenteral administration are solutions, preferably oil-based or aqueous solutions, furthermore suspensions, emulsions or implants, and suitable for topical application are ointments, creams or powders or also as nasal sprays. The novel compounds may also be lyophilised and the resultant
35 lyophilisates used, for example, to prepare injection preparations. The preparations indicated may be sterilised and/or comprise assistants, such

as lubricants, preservatives, stabilisers and/or wetting agents, emulsifying agents, salts for modifying the osmotic pressure, buffer substances, colorants and flavours and/or a plurality of further active ingredients, for example one or more vitamins.

5

The compounds of the formula I and their physiologically acceptable salts can be used for combating and preventing thromboembolic disorders, such as thrombosis, myocardial infarction, arteriosclerosis, inflammation, 10 apoplexia, angina pectoris, restenosis after angioplasty, claudicatio intermittens, tumours, tumour diseases and/or tumour metastases.

In general, the substances according to the invention are preferably 15 administered in doses between about 1 and 500 mg, in particular between 5 and 100 mg, per dosage unit. The daily dose is preferably between about 0.02 and 10 mg/kg of body weight. However, the specific dose for each patient depends on a wide variety of factors, for example on the 20 efficacy of the specific compound employed, on the age, body weight, general state of health, sex, on the diet, on the time and method of administration, on the excretion rate, medicament combination and severity of the particular illness to which the therapy applies. Oral 25 administration is preferred.

The invention furthermore relates to medicaments comprising at least one compound of the formula I and/or its pharmaceutically usable derivatives, solvates and stereoisomers, including mixtures thereof in all ratios, and at 30 least one further medicament active ingredient.

The invention also relates to a set (kit) consisting of separate packs of

35

- (a) an effective amount of a compound of the formula I and/or its pharmaceutically usable derivatives, solvates and stereoisomers, including mixtures thereof in all ratios,
and
5 (b) an effective amount of a further medicament active ingredient.

The set comprises suitable containers, such as boxes, individual bottles, bags or ampoules. The set may, for example, comprise separate ampoules
10 each containing an effective amount of a compound of the formula I and/or its pharmaceutically usable derivatives, solvates and stereoisomers, including mixtures thereof in all ratios,
and an effective amount of a further medicament active ingredient in
15 dissolved or lyophilised form.

The invention furthermore relates to the use of compounds of the formula I and/or their pharmaceutically usable derivatives, solvates and stereoisomers, including mixtures thereof in all ratios,
20 for the preparation of a medicament for the treatment of thrombosis, myocardial infarction, arteriosclerosis, inflammation, apoplexia, angina pectoris, restenosis after angioplasty, claudicatio intermittens, tumours, tumour diseases and/or tumour metastases,
25 in combination with at least one further medicament active ingredient.

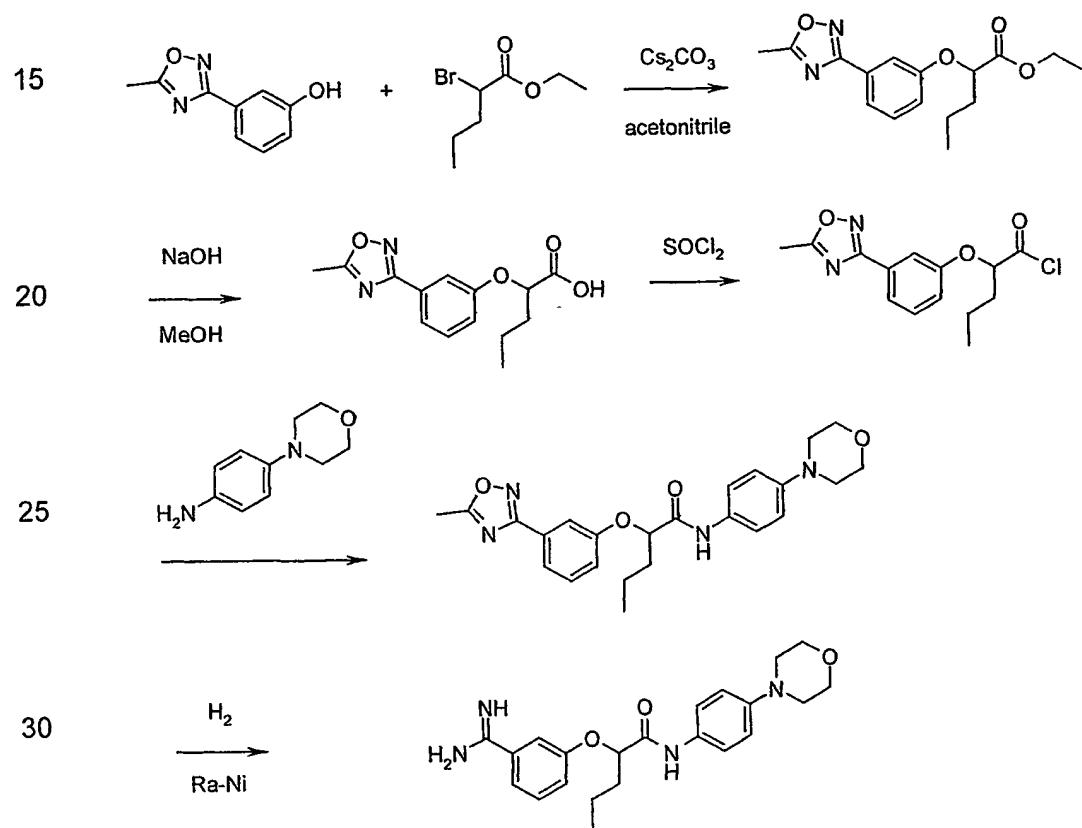
Above and below, all temperatures are given in °C. In the following examples, "conventional work-up" means that water is added if necessary, the pH is adjusted, if necessary, to between 2 and 10, depending on the constitution of the end product, the mixture is extracted with ethyl acetate or dichloromethane, the phases are separated, the organic phase is dried over sodium sulfate and evaporated, and the product is purified by
30
35

chromatography on silica gel and/or by crystallisation. Rf values on silica gel; eluent: ethyl acetate/methanol 9:1.

Mass spectrometry (MS): EI (electron impact ionisation) M^+
FAB (fast atom bombardment) $(M+H)^+$
ESI (electrospray ionisation) $(M+H)^+$

Example 1

10 2-(3-amidinophenoxy)-*N*-(4-morpholin-4-ylphenyl)valeramide is prepared as indicated in the following scheme:



35 1. 380 g (1.17 mol) of caesium carbonate are added to a solution of 200 g (1.135 mol) of 3-(5-methyl-1,2,4-oxadiazol-3-yl)phenol in 2.5 l of

acetonitrile. A solution of 190 ml (1.15 mol) of ethyl 2-bromoalate in 0.5 l of acetonitrile is subsequently added. The reaction mixture is stirred at room temperature for 24 hours and then filtered, and the filtrate is evaporated, giving ethyl 2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)-
5 phenoxy]pentanoate as a yellowish oil; ESI 305.

2. 2.1 l of 1N sodium hydroxide solution are added to a solution of 347 g (1.14 mol) of ethyl 2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]-
10 pentanoate in 1.4 l of methanol, and the mixture is stirred at room temperature for 24 hours. The reaction mixture is diluted with 1 l of water and extracted twice with 1 l of tert-butyl methyl ether each time. The aqueous phase is acidified to a pH of 2 using concentrated hydrochloric
15 acid and extracted twice with 2 l of tert-butyl methyl ether each time. The combined organic phases are evaporated, and the residue is recrystallised from 100 ml of toluene: 2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]pentanoic acid is obtained as colourless crystals of m.p. 98-
20 100°.

3. 3.50 g (12.7 mmol) of 2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)-
phenoxy]pentanoic acid are dissolved in 10 ml of thionyl chloride, and
25 the mixture is heated at the boil for 2 hours. The reaction mixture is evaporated, the residue is taken up in toluene, and the solution is re-evaporated, giving 2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]-
pentanoyl chloride as a yellowish solid.

30 4. 200 mg of 4-dimethylaminopyridine on polystyrene are added to a solution of 100 mg (0.339 mmol) of 2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)-
phenoxy]pentanoyl chloride and 60.4 mg (0.339 mmol) of 4-morpholin-
35 4-ylaniline in 6 ml of dichloromethane, and the mixture is stirred at room temperature for 18 hours. The reaction mixture is filtered, and the filtrate

is evaporated, giving N-(4-morpholin-4-ylphenyl)-2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]pentanoamide as a colourless solid, ESI 437.

5 200 mg of water-moist Raney nickel and 0.25 ml of acetic acid are added to a solution of 140 mg (0.321 mmol) of N-(4-morpholin-4-ylphenyl)-2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]pentanoamide in 6 ml of methanol, and the mixture is hydrogenated at room temperature and atmospheric pressure for 18 hours. The reaction mixture is filtered, 10 and the residue is evaporated, giving 2-(3-amidinophenoxy)-N-(4-morpholin-4-ylphenyl)valeramide, acetate, ESI 397;
IC₅₀ (Xa) = 3 x 10⁻⁷ M;
IC₅₀ (VIIa) = 4.9 x 10⁻⁷ M.

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The following compound is obtained analogously:

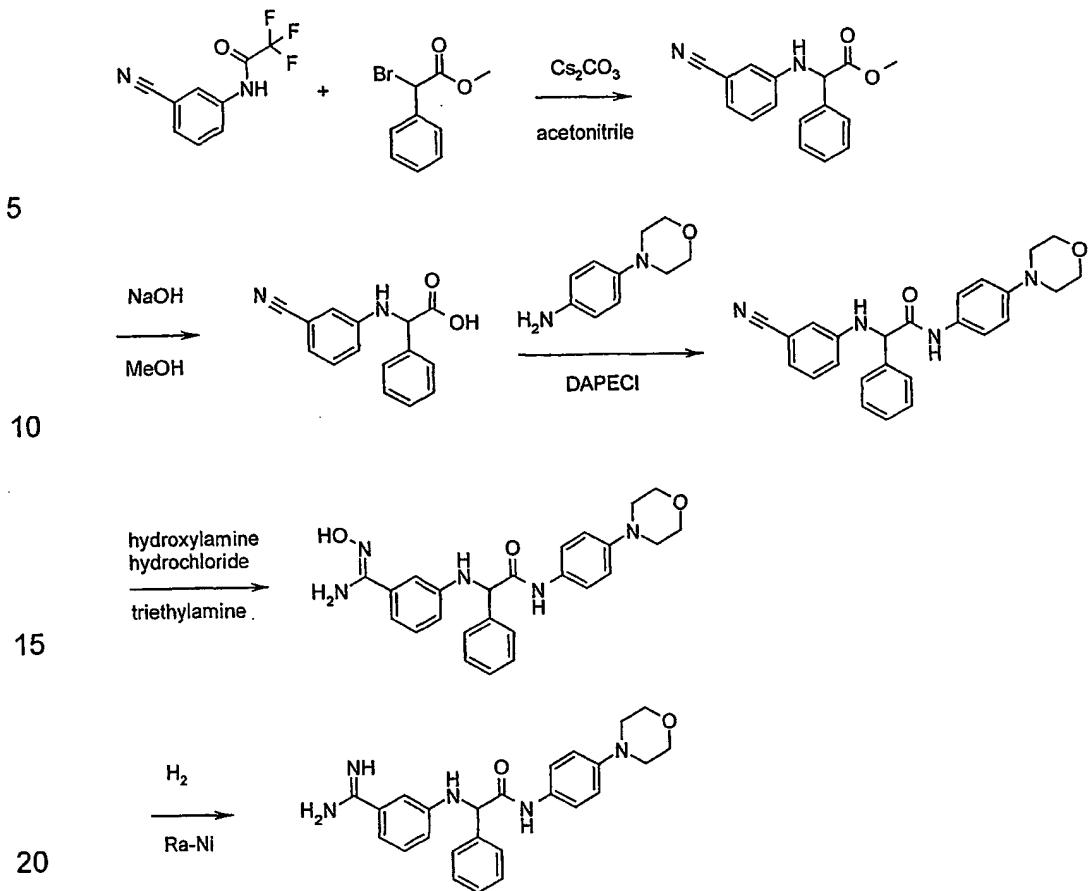
20 2-(3-amidinophenoxy)-N-(4-morpholin-4-ylphenyl)-2-phenylacetamide, diacetate, ESI 431; IC₅₀ (Xa) = 9.0 x 10⁻⁸ M;
IC₅₀ (VIIa) = 6.0 x 10⁻⁸ M.

Example 2

25 2-(3-amidinophenylamino)-N-(4-morpholin-4-ylphenyl)-2-phenylacetamide is prepared as indicated in the following scheme:

30

35



1. 14.3 g (44.0 mmol) of caesium carbonate are added to a solution of 8.6 g (40.2 mmol) of *N*-(3-cyanophenyl)[2.2.2]trifluoroacetamide in 200 ml of acetonitrile. 7.2 ml (44.4 mmol) of methyl 2-bromo-2-phenylacetate are subsequently added. The reaction mixture is heated at the boil with stirring for 3 hours, then filtered, the filtrate is evaporated, and the residue is chromatographed on a silica-gel column with dichloromethane/petroleum ether, giving methyl (3-cyanophenylamino)-2-phenylacetate as a colourless oil; ESI 267.

2. 50 ml of 1N sodium hydroxide solution are added to a solution of 9.5 g (35.6 mmol) of methyl (3-cyanophenylamino)-2-phenylacetate in 150 ml of methanol, and the mixture is stirred at room temperature for

21 hours. The reaction mixture is evaporated, the residue is acidified to pH 1 using 25% HCl, and the resultant precipitate is filtered off, giving (3-cyanophenylamino)-2-phenylacetic acid as colourless crystals of m.p. 159-162°.

5

3. 87 µl (0.793 mmol) of 4-methylmorpholine are added to a solution of 200 mg (0.793 mmol) of (3-cyanophenylamino)-2-phenylacetic acid, 141 mg (0.793 mmol) of 4-morpholin-4-ylaniline, 152 mg (0.793 mmol) 10 of N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide hydrochloride (DAPECI) and 121 mg (0.793 mmol) of hydroxybenzotriazole hydrate (HOBr) in 3 ml of DMF, and the mixture is stirred at room temperature for 48 hours. The reaction mixture is introduced into water, and the 15 precipitate is filtered off, giving 2-(3-cyanophenylamino)-N-(4-morpholin-4-ylphenyl)-2-phenylacetamide as a colourless solid; ESI 413.

4. 149 mg (2.15 mmol) of hydroxylamine hydrochloride, 0.30 ml 20 (2.15 mmol) of triethylamine and a few grains of 0.3 nm molecular sieve are added to a solution of 295 mg (0.715 mmol) of 2-(3-cyanophenylamino)-N-(4-morpholin-4-ylphenyl)-2-phenylacetamide in 5 ml of methanol, and the mixture is heated at 70°C for 18 hours. The reaction mixture is filtered, the filtrate is evaporated, and the residue is taken up in 25 water. The resultant precipitate is filtered off, giving 2-[3-(N-hydroxyamidino)phenylamino]-N-(4-morpholin-4-ylphenyl)-2-phenylacetamide as a colourless solid; ESI 446.

30

5. 500 mg of water-moist Raney nickel and 0.1 ml of acetic acid are added to a solution of 150 mg (0.337 mmol) of 2-[3-(N-hydroxyamidino)phenylamino]-N-(4-morpholin-4-ylphenyl)-2-phenylacetamide in 35 5 ml of methanol, and the mixture is hydrogenated at room temperature and atmospheric pressure for 18 hours. The reaction mixture is filtered,

and the residue is evaporated, giving 2-(3-amidinophenylamino)-N-(4-morpholin-4-ylphenyl)-2-phenylacetamide, triacetate, ESI 430;
IC₅₀ (Xa) = 3.4 x 10⁻⁸ M;
IC₅₀ (VIIa) = 2.2 x 10⁻⁸ M;

5

The following compound is obtained analogously:

10 2-(3-amidinophenylamino)-N-(4-dimethylaminophenyl)-4-methylvaler-
amide, diacetate, ESI 368; IC₅₀ (Xa) = 1.6 x 10⁻⁷ M;
IC₅₀ (VIIa) = 2.1 x 10⁻⁸ M.

15 Example 3

15

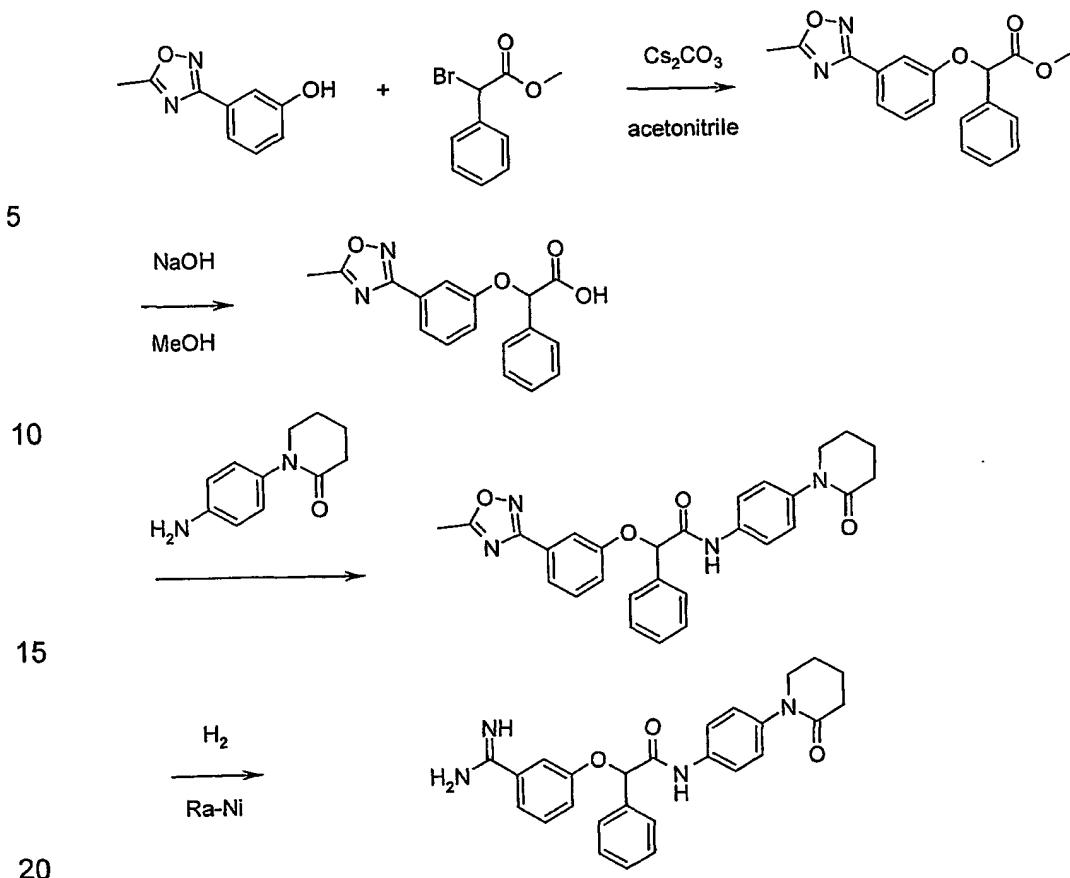
20 2-(3-amidinophenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacet-
amide is prepared as indicated in the following scheme:

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- 40 -



1. 11.5 g (33.5 mmol) of caesium carbonate and 7.67 g (35.2 mmol) of methyl 2-bromo-2-phenylacetate are added to a solution of 5.90 g (33.5 mmol) of 3-(5-methyl-1,2,4-oxadiazol-3-yl)phenol in 50 ml of acetonitrile. The reaction mixture is stirred at room temperature for 24 hours, then filtered, and the filtrate is evaporated, giving methyl [3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]-2-phenylacetate as a yellowish oil; ESI 325.
 2. 30 ml of 1N sodium hydroxide solution are added to a solution of 10.1 g (31.0 mmol) of methyl [3-(5-methyl-1,2,4-oxadiazol-3-yl)-phenoxy]-2-phenylacetate in 30 ml of methanol, and the mixture is heated at 80°C for 3 hours. The reaction mixture is evaporated, 1N HCl is added, and the mixture is extracted with ethyl acetate. The organic
- 25
- 30
- 35

phase is dried and evaporated, and the residue is recrystallised from ether, giving [3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]-2-phenylacetic acid as colourless crystals; ESI 311.

- 5 3. 35 µl (0.322 mmol) of 4-methylmorpholine are added to a solution of 100 mg (0.322 mmol) of [3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]-2-phenylacetic acid, 61.3 mg (0.322 mmol) of 1-(4-aminophenyl)piperidin-2-one (prepared from 1-(4-nitrophenyl)piperidin-2-one by hydrogenation using Raney nickel as catalyst), 61.7 mg (0.322 mmol) of *N*-(3-dimethylaminopropyl)-*N'*-ethylcarbodiimide hydrochloride (DAPECI) and 49.3 mg (0.322 mmol) of hydroxybenzotriazole hydrate (HOBr) in 3 ml of DMF, and the mixture is stirred at room temperature for 48 hours. The reaction mixture is introduced into aqueous sodium carbonate solution, and the precipitate is filtered off, giving 2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]-*N*-(4-(2-oxopiperidin-1-yl)phenyl)-2-phenylacetamide as a colourless solid; ESI 483.
- 10 15 20 25 30
4. 200 mg of water-moist Raney nickel and 0.2 ml of acetic acid are added to a solution of 122 mg (0.253 mmol) of 2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]-*N*-(4-(2-oxopiperidin-1-yl)phenyl)-2-phenylacetamide in 5 ml of methanol, and the mixture is hydrogenated at room temperature and atmospheric pressure for 18 hours. The reaction mixture is filtered, and the residue is evaporated, giving 2-(3-amidinophenoxy)-*N*-(4-(2-oxopiperidin-1-yl)phenyl)-2-phenylacetamide, acetate (EMD 388547), ESI 443; IC₅₀ (Xa) = 3.2 × 10⁻⁸ M; IC₅₀ (VIIa) = 1.4 × 10⁻⁸ M.

The following compounds are obtained analogously:

2-(3-amidinophenoxy)-N-[4-(2-oxopyrrolidin-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 429; IC₅₀ (Xa) = 6.5 x 10⁻⁸ M;
IC₅₀ (VIIa) = 3.9 x 10⁻⁸ M;

5

2-(3-amidinophenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]valeramide,
acetate, ESI 409; IC₅₀ (Xa) = 7.0 x 10⁻⁸ M;
IC₅₀ (VIIa) = 3.3 x 10⁻⁸ M;

10

2-(3-amidinophenoxy)-N-[4-(2-oxopyrrolidin-1-yl)phenyl]valeramide,
acetate, ESI 395; IC₅₀ (Xa) = 1.3 x 10⁻⁷ M;
IC₅₀ (VIIa) = 8.3 x 10⁻⁸ M;

15

2-(3-amidinophenoxy)-N-[4-(2-oxo-1*H*-pyridin-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-N-[4-(3-oxomorpholin-4-yl)phenyl]-2-phenylacetamide,

20

2-(3-amidinophenoxy)-N-[4-(4-oxo-1*H*-pyridin-1-yl)phenyl]-2-phenylacetamide,
2-(3-(N-Hydroxyamidino)phenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,

25

2-(3-amidinophenoxy)-N-[4-(2,6-dioxopiperidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-N-[4-(3-oxo-2*H*-piperazin-1-yl)phenyl]-2-phenylacetamide,

30

2-(3-amidinophenoxy)-N-[2-fluoro-4-(2-oxo-1*H*-pyridin-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-N-[4-(2,5-dioxopyrrolidin-1-yl)phenyl]-2-phenylacetamide,

35

2-(3-amidinophenoxy)-N-[5-(2-oxopiperidin-1-yl)pyridin-2-yl]-2-phenylacetamide,

- 2-(3-amidinophenoxy)-N-[4-(2-oxo-1,3-oxazolidin-3-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-N-[4-(2-oxo-1*H*-pyridin-1-yl)benzyl]-2-phenylacetamide,
5 2-(3-amidinophenoxy)-N-[4-(2-oxopiperidin-1-yl)benzyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-N-[4-(2-oxopiperidin-1-yl)butyl]-2-phenylacetamide,
10 2-(3-amidinophenoxy)-N-[4-(3-oxo-2*H*-pyridazin-2-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-N-[2-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
15 2-(3-amidinophenoxy)-N-(4-dimethylaminobenzyl)-2-phenylacetamide, ESI 403;
2-(3-amidinophenoxy)-N-[3-(morpholin-4-yl)propyl]-2-phenylacetamide,
20 2-(3-amidinophenoxy)-N-[3-(piperidin-1-yl)propyl]-2-phenylacetamide,
ESI 381;
2-(3-amidinophenoxy)-N-(4-dimethylaminophenyl)-2-phenylacetamide, ESI 389.
25 Analogous reaction of [3-(5-methyl-1,2,4-oxadiazol-3-yl)phenylamino]-2-phenylacetic acid and
1-(4-aminophenyl)piperidin-2-one,
30 2-(morpholin-4-yl)ethylamine,
4-dimethylaminobenzylamine,
3-(morpholin-4-yl)propylamine,
3-(piperidin-1-yl)propylamine,
35 and subsequent hydrogenation gives

2-(3-amidinophenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,

5 2-(3-amidinophenylamino)-*N*-[2-(morpholin-4-yl)ethyl]-2-phenylacetamide,

10 2-(3-amidinophenylamino)-*N*-(4-dimethylaminobenzyl)-2-phenylacetamide,

15 2-(3-amidinophenylamino)-*N*-[3-(morpholin-4-yl)propyl]-2-phenylacetamide,

20 2-(3-amidinophenylamino)-*N*-[3-(piperidin-1-yl)propyl]-2-phenylacetamide.

25 Analogous reaction of [3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]-2-valeric acid and

2-(morpholin-4-yl)ethylamine,

4-dimethylaminobenzylamine,

20 3-(morpholin-4-yl)propylamine,

3-(piperidin-1-yl)propylamine,

and subsequent hydrogenation gives

25 2-(3-amidinophenoxy)-*N*-[2-(morpholin-4-yl)ethyl]valeramide,
ESI 349;

2-(3-amidinophenoxy)-*N*-(4-dimethylaminobenzyl)valeramide,

30 ESI 369;
2-(3-amidinophenoxy)-*N*-[3-(morpholin-4-yl)propyl]valeramide,

ESI 363;
2-(3-amidinophenoxy)-*N*-[3-(piperidin-1-yl)propyl]valeramide,

35 ESI 347.

Example 4

Analogously to Examples 1-3, reaction of (3-cyanophenoxy)-2-phenyl-acetic acid and

- 5 1-(4-aminophenyl)piperidin-2-one,
 2-(morpholin-4-yl)ethylamine,
 4-dimethylaminobenzylamine,
 3-(morpholin-4-yl)propylamine,
10 3-(piperidin-1-yl)propylamine,

followed by hydrolysis using aqueous sodium hydroxide solution gives

- 15 2-(3-aminocarbonylphenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, ESI 444;
 2-(3-aminocarbonylphenoxy)-N-[2-(morpholin-4-yl)ethyl]-2-phenylacetamide,
20 2-(3-aminocarbonylphenoxy)-N-(4-dimethylaminobenzyl)-2-phenylacetamide,
 2-(3-aminocarbonylphenoxy)-N-[3-(morpholin-4-yl)propyl]-2-phenylacetamide,
25 2-(3-aminocarbonylphenoxy)-N-[3-(piperidin-1-yl)propyl]-2-phenylacetamide.

Example 5

- 30 The following compounds are obtained analogously to Example 1:

- 35 2-(3-amidinophenoxy)-N-(4-morpholin-4-ylbenzyl)valeramide,
 diacetate, ESI 411;

- 2-(3-amidinophenoxy)-N-(4-morpholin-4-ylbenzyl)-2-phenylacetamide, diacetate, ESI 445;
- 2-(3-amidinophenoxy)-N-[4-(3-oxomorpholin-4-yl)phenyl]-2-phenylacetamide, acetate, ESI 445;
- 5 2-(3-amidinophenoxy)-N-[3-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 443;
- 10 2-(3-amidinophenoxy)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 457;
- 15 (2R)-2-(3-amidinophenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]valeramide, acetate, ESI 409;
- 2-(3-amidinophenoxy)-N-[2-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 461;
- 20 2-(3-amidinophenoxy)-N-[2-fluoro-4-(2-caprolactam-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 475;
- 2-(3-amidinophenoxy)-N-[3-methyl-4-(2-caprolactam-1-yl)phenyl]-valeramide, acetate, ESI 437;
- 25 2-(3-amidinophenoxy)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-valeramide, acetate, ESI 423;
- 2-(3-amidinophenoxy)-N-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-valeramide, acetate, ESI 427;
- 30 2-(3-amidinophenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]acetamide, acetate, ESI 367;
- 2-(2-amidinophenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 443;
- 35 2-(4-amidinophenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 443;
- 3-(3-amidinophenyl)-N-[4-(2-oxopiperidin-1-yl)phenyl]propionamide, ESI 365;
- 2-(3-amidinophenoxy)-N-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, ESI 461; ~

5 3-(3-amidinophenyl)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-propionamide, ESI 379;

10 (2*S*)-2-(3-amidinophenoxy)-*N*-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-valeramide, acetate, ESI 427;

15 2-(3-amidinophenoxy)-*N*-[3-methyl-4-(2-caprolactam-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 471;

20 2-(3-amidinophenylmethyl)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]valeramide, acetate, ESI 407;

25 2-(3-amidinophenylmethyl)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]valeramide, acetate, ESI 421;

30 2-(3-amidinophenoxy)-*N*-[4-(2-oxopiperazin-1-yl)phenyl]valeramide, diacetate, ESI 410;

35 (2*S*)-2-(3-amidinophenoxy)-*N*-[4-(2-caprolactam-1-yl)phenyl]valeramide, acetate, ESI 423;

40 (2*S*)-2-(3-amidinophenoxy)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]valeramide, acetate, ESI 423;

45 2-(3-amidinophenoxy)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide, acetate, ESI 437;

50 2-(3-amidinophenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide, acetate, ESI 423;

55 (2*R*)-2-(3-amidinophenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]valeramide, acetate, ESI 409;

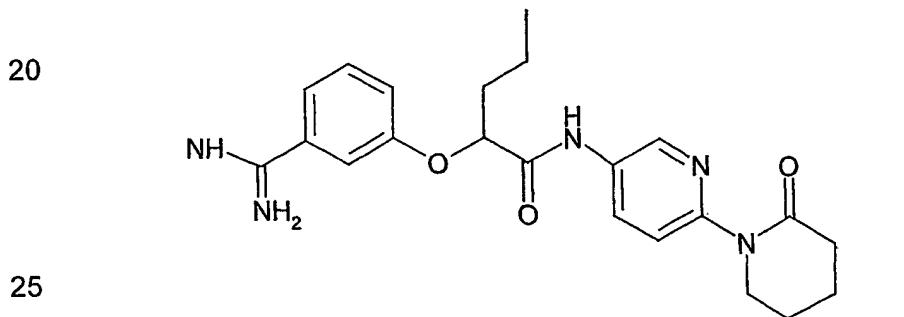
60 2-(3-amidinophenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]propionamide, acetate, ESI 381;

65 2-(3-amidinophenoxy)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-propionamide, acetate, ESI 395;

70 2-(3-amidinophenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]butyramide, acetate, ESI 395;

75 2-(3-amidinophenoxy)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-butyramide, acetate, ESI 409;

- (2S)-2-(3-amidinophenoxy)-N-[3-methyl-4-(2-caprolactam-1-yl)-phenyl]valeramide, acetate, ESI 437;
- 2-(3-amidinophenoxy)-N-[4-(2-oxopyridin-1-yl)phenyl]valeramide, acetate, ESI 405;
- 5 2-(3-amidinophenoxy)-N-[2,5-dimethyl-4-(2-oxopiperidin-1-yl)phenyl]-valeramide, acetate, ESI 437;
- 2-(3-amidinophenoxy)-N-[3-methyl-4-(2-oxopyrrolidin-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 443;
- 10 2-(3-amidinophenoxy)-N-[3-methyl-4-(2-oxopyrrolidin-1-yl)phenyl]-valeramide, acetate, ESI 409;
- 2-(3-amidinophenoxy)-N-[2-methyl-4-(2-oxopiperidin-1-yl)phenyl]-valeramide, acetate, ESI 423;
- 15 2-(3-amidinophenoxy)-N-(2-oxo-3,4,5,6-tetrahydro-2H-1,2'-bipyridinyl-5'-yl)valeramide, diacetate, ESI 410



- 30 2-(3-amidinophenoxy)-N-[2-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 457;
- 2-(3-amidinophenoxy)-N-(2-oxo-3,4,5,6-tetrahydro-2H-1,2'-bipyridinyl-5'-yl)-2-phenylacetamide, diacetate, ESI 444;
- 35 2-[(3-N-methoxycarbonylamidino)phenoxy]-N-[4-(2-oxopiperidin-1-yl)phenyl]valeramide, ESI 467;

- 2-(3-amidinophenoxy)-*N*-[3-fluoro-4-(2-oxopyrrolidin-1-yl)phenyl]-valeramide, acetate, ESI 413;
- 2-(3-amidinophenoxy)-*N*-[3-fluoro-4-(2-oxopyrrolidin-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 447;
- 5 2-(3-amidinophenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]caproamide, acetate, ESI 423;
- 10 2-(3-amidinophenoxy)-*N*-[4-(2-oxopyrrolidin-1-yl)phenyl]butyramide, acetate, ESI 381;
- 15 2-(3-amidinophenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-3-methylbutyramide, acetate, ESI 409;
- 20 2-(3-amidinophenoxy)-*N*-[4-ethyl-3-(2-oxopyrrolidin-1-yl)phenyl]valeramide, acetate, ESI 423;
- 25 2-(3-amidinophenoxy)-*N*-[3-fluoro-4-(2-caprolactam-1-yl)phenyl]valeramide, acetate, ESI 441;
- 30 2-(3-amidinophenoxy)-*N*-[3-fluoro-4-(2-caprolactam-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 475;
- 35 2-(3-amidinophenoxy)-*N*-[4-(2,6-dioxopiperidin-1-yl)phenyl]valeramide, acetate, ESI 423;
- 3-(3-amidinophenyl)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylpropionamide, acetate, ESI 441;
- 2-(3-amidinophenyl)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylpropionamide, acetate, ESI 455;
- 2-(3-amidinophenoxy)-*N*-[3-methoxy-4-(2,5-dioxopyrrolidin-1-yl)phenyl]valeramide, acetate, ESI 439;
- 30 2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]-*N*-[4-(2,6-dioxopiperidin-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 497;
- 2-(3-amidinophenoxy)-*N*-[4-(2,6-dioxopiperidin-1-yl)phenyl]-2-phenylacetamide, ESI 457;
- 35 2-(3-amidinophenoxy)-*N*-[3-trifluoromethyl-4-(2,6-dioxopiperidin-1-yl)phenyl]valeramide, ESI 491;

2-(3-amidinophenoxy)-N-[3-chloro-4-(2,5-dioxopyrrolidin-1-yl)phenyl]-valeramide, acetate, ESI 443;

2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]-N-[3-trifluoromethyl-4-(2,5-dioxopyrrolidin-1-yl)phenyl]-2-valeramide, ESI 517;

5 2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]-N-[3-chloro-4-(2,5-dioxopyrrolidin-1-yl)phenyl]-2-valeramide, ESI 483;

10 2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]-N-[3-methoxy-4-(2,5-dioxopyrrolidin-1-yl)phenyl]-2-valeramide, ESI 479;

15 2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]-N-[4-(2,5-dioxopyrrolidin-1-yl)phenyl]-2-valeramide, ESI 449;

3-(3-amidinophenoxy)-N-[3-chloro-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 477;

20 3-(3-amidinophenoxy)-N-[3-chloro-4-(2-oxopiperidin-1-yl)phenyl]-valeramide, acetate, ESI 443;

3-(3-amidinophenoxy)-N-[3-chloro-4-(2-oxopiperidin-1-yl)phenyl]-butyramide, acetate, ESI 429.

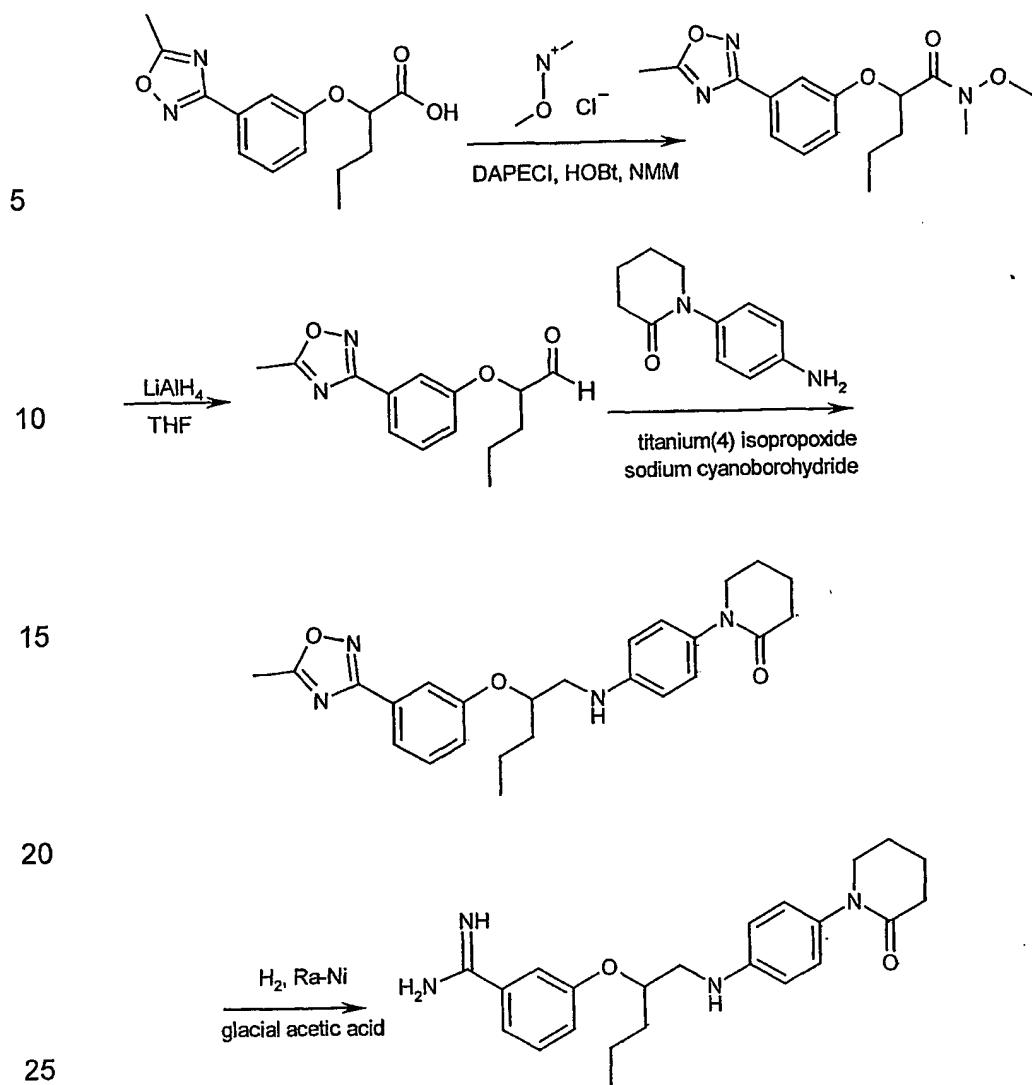
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Example 5a

25 1-{4-[2-(3-Amidinophenoxy)pentylamino]phenyl}piperidin-2-one, diacetate, ESI 395, is prepared in accordance with the following scheme:

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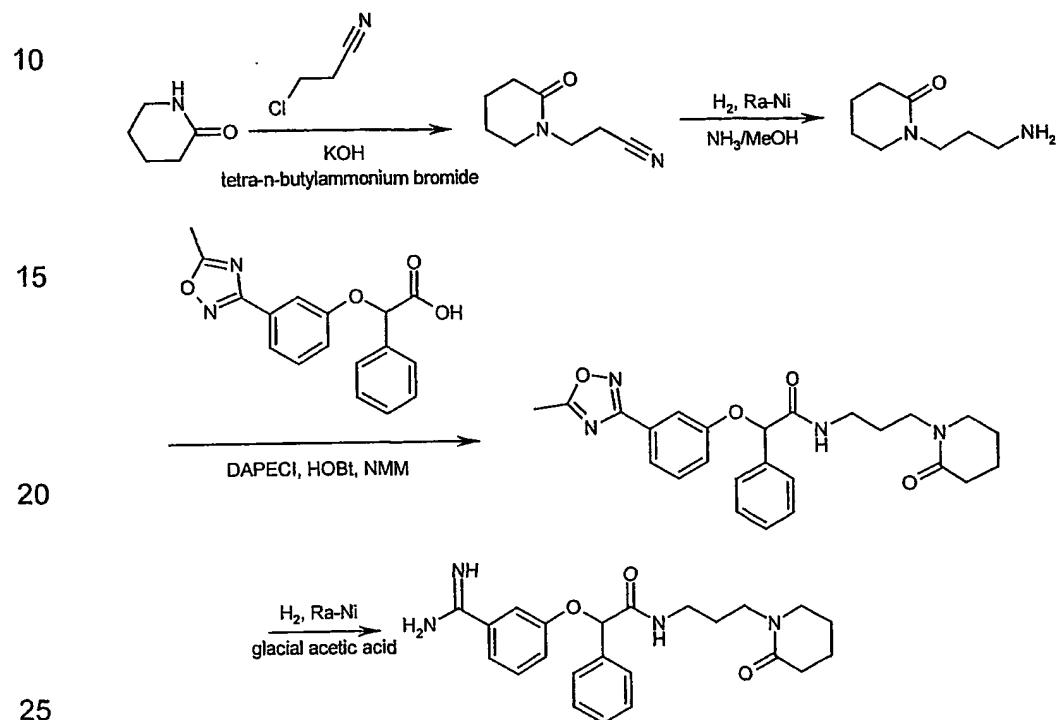


The following compounds are obtained analogously:

30 1-{4-[2-(3-amidinophenoxy)pentylamino]-2-methylphenyl}piperidin-2-one,
diacetate, ESI 409;
1-{4-[2-(3-amidinophenoxy)-2-phenylethoxy]phenyl}piperidin-2-one,
acetate, ESI 430.

Example 5b

5 2-(3-Amidinophenoxy)-N-[3-(2-oxopiperidin-1-yl)propyl]-2-phenylacetamide, acetate, ESI 409, is prepared in accordance with the following scheme:

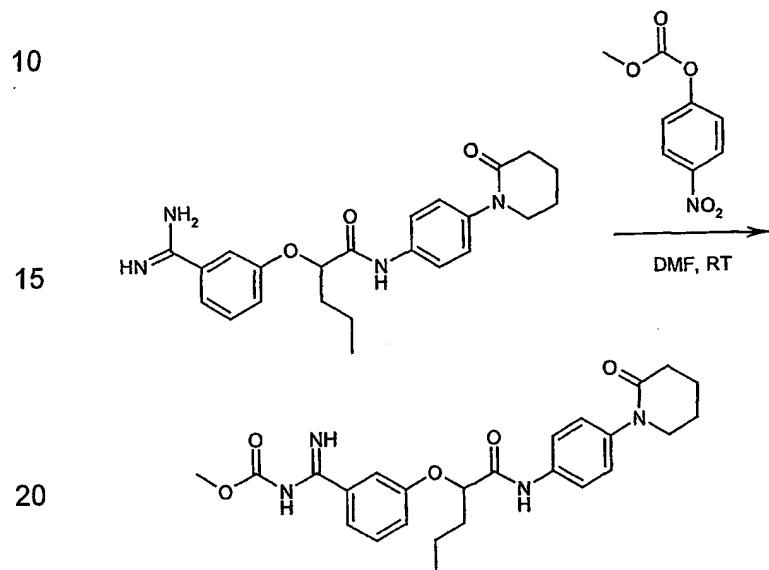


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Example 5c

5 2-[(3-N-Ethoxycarbonylamidino)phenoxy]-N-[4-(2-oxopiperidin-1-yl)phenyl]-
valeramide, ESI 481, is prepared in accordance with the following scheme:



25 The following compounds are obtained analogously:

2-[(3-N-methoxycarbonylamidino)phenoxy]-N-[4-(2-oxopiperidin-1-yl)]-
phenyl]-2-phenylacetamide, ESI 501;
2-[(3-N-ethoxycarbonylamidino)phenoxy]-N-[4-(2-oxopiperidin-1-yl)]-
phenyl]-2-phenylacetamide, ESI 515.

Example 6

5 The following compounds are obtained analogously to Example 2:

2-(3-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide, diacetate, ESI 422;

10 2-(3-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, diacetate, ESI 442;

 2-(3-amidinophenylamino)-N-[4-(2-oxopyrrolidin-1-yl)phenyl]-4-methylvaleramide, diacetate, ESI 408;

15 2-(3-amidinophenylamino)-N-[4-(2-oxopyrrolidin-1-yl)phenyl]-2-phenylacetamide, diacetate, ESI 428;

 2-(3-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide, acetate, ESI 460;

20 2-(3-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]acetamide, acetate, ESI 366;

 (2S)-2-(3-amidinophenylamino)-N-[4-(2-caprolactam-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 456;

25 (2S)-2-(3-amidinophenylamino)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 456;

 (2S)-2-(3-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, diacetate, ESI 442;

30 2-(3-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]caproamide, acetate, ESI 422;

 2-(3-amidinophenylamino)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide, diacetate, ESI 474;

35 (2S)-2-(2-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 442;

- (2R)-2-(2-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 442;
- (2S)-2-(3-amidinophenylamino)-N-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, ESI 460;
- 5 2-(3-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-(3-fluorophenyl)acetamide, acetate, ESI 460;
- 2-(3-amidinophenylamino)-N-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-2-(3-fluorophenyl)acetamide, acetate, ESI 478;
- 10 (2R)-2-(3-amidinophenylamino)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide, ESI 436;
- 2-(3-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]valeramide, diacetate, ESI 408;
- 15 (2R)-2-(3-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide, acetate, ESI 422;
- (2R)-2-(3-amidinophenylamino)-N-[4-(2-caprolactam-1-yl)phenyl]-4-methylvaleramide, acetate, ESI 436;
- 20 2-(3-amidinophenylamino)-N-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide, acetate, ESI 478;
- 2-(3-amidinophenylamino)-N-[3-methyl-4-(2-caprolactam-1-yl)phenyl]-2-(2-fluorophenyl)acetamide, acetate, ESI 488;
- 25 (2R)-2-(3-amidinophenylamino)-N-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide, ESI 440;
- 2-(3-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-4-trifluoro-3-methylbutyramide, acetate, ESI 462;
- 30 (2S)-2-(3-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)benzyl]-2-phenylacetamide, acetate, ESI 456;
- (2S)-2-(3-amidinophenylamino)-N-[4-(3-oxo-2-azabicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide, acetate, ESI 468;
- 35 2-(3-amidinophenylamino)-N-[3-trifluoromethyl-4-(3-oxo-2-azabicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide, ESI 536;

2-(3-amidino-4-fluorophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-
2-phenylacetamide, diacetate, ESI 460;

5

Example 6a

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2-(3-Amidinophenylamino)-N-methyl-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-
phenylacetamide, acetate, ESI 456, is prepared in accordance with the
following scheme:

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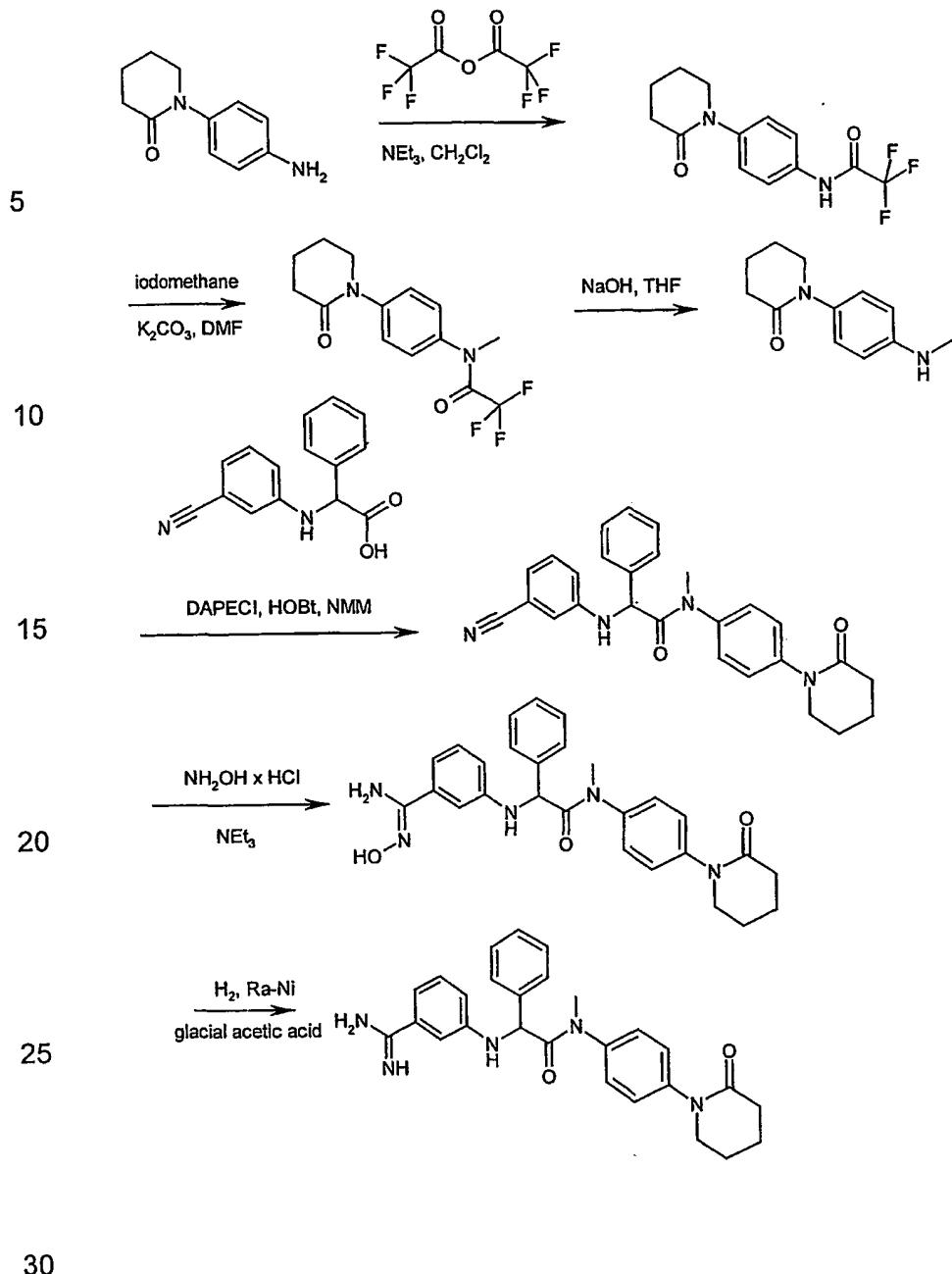
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Example 7

5 The following are obtained analogously to Example 4:

- 2-(3-aminocarbonylphenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]valer-
amide, ESI 410;
- 10 2-(3-aminocarbonylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-
phenylacetamide, ESI 443;
- 15 2-(3-aminocarbonylphenoxy)-N-[4-(2-oxo-1*H*-pyridin-1-yl)benzyl]-2-
phenylacetamide, ESI 454;
- 20 2-(3-aminocarbonylphenoxy)-N-[4-(2-caprolactam-1-yl)phenyl]-2-
phenylacetamide, ESI 457;
- 25 2-(3-aminocarbonylphenoxy)-N-[4-(2-caprolactam-1-yl)phenyl]-2-
(2-fluorophenyl)acetamide, ESI 461;
- 30 3-(3-aminocarbonylphenyl)-N-[4-(2-oxopiperidin-1-yl)phenyl]propion-
amide, ESI 366;
- 35 2-(3-aminocarbonylphenoxy)-N-[4-(2-oxo-1*H*-pyridin-1-yl)phenyl]-2-
phenylacetamide, ESI 440;
- 2-(3-aminocarbonylphenoxy)-N-[3-methyl-4-(2-oxopiperidin-1-yl)-
phenyl]-2-phenylacetamide, ESI 458;
- 2-(3-aminocarbonylphenoxy)-N-[4-(4-oxo-4*H*-pyridin-1-yl)phenyl]-2-
phenylacetamide, ESI 458;
- 2-(3-aminocarbonylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-
acetamide, ESI 367;
- (2S)-2-(3-aminocarbonylphenylamino)-N-[3-methyl-4-(2-oxopiperidin-
1-yl)phenyl]-2-phenylacetamide, ESI 457;

- (2S)-2-(3-aminocarbonylphenylamino)-N-[4-(2-caprolactam-1-yl)-phenyl]-2-phenylacetamide, ESI 457;
- (2R)-2-(3-aminocarbonylphenylamino)-N-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide, ESI 443;
- 5 (2S)-2-(3-aminocarbonylphenylamino)-N-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide, ESI 443;
- 2-(3-aminocarbonylphenylamino)-N-[4-(2-caprolactam-1-yl)phenyl]-2-(2-fluorophenyl)acetamide, ESI 475;
- 10 2-(3-aminocarbonylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-(2-thienyl)acetamide, ESI 449;
- 2-(4-aminocarbonylphenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, ESI 444;
- 15 2-(2-aminocarbonylphenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, ESI 444;
- 2-(3-aminocarbonylphenoxy)-N-[2-fluoro-4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide, ESI 462;
- 20 2-(3-aminocarbonylphenoxy)-N-[2-fluoro-4-(2-caprolactam-1-yl)-phenyl]-2-phenylacetamide, ESI 475;
- (2S)-2-(2-aminocarbonylphenyl)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, ESI 443;
- 25 (2R)-2-(2-aminocarbonylphenyl)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, ESI 443;
- 2-(3-aminocarbonylphenoxy)-N-[3-methyl-4-(2-caprolactam-1-yl)-phenyl]-2-phenylacetamide, ESI 472;
- 30 (2S)-2-(3-aminocarbonylphenylamino)-N-[4-(2-oxopyridin-1-yl)-phenyl]-2-phenylacetamide, ESI 439;
- 2-(3-aminocarbonylphenylamino)-N-[4-(2-oxopiperidin-1-yl)-phenyl]valeramide, ESI 409;
- 35 2-(3-aminocarbonyl-4-fluorophenylamino)-N-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide, ESI 461;

2-(3-aminocarbonylphenylamino)-N-[3-trifluoromethyl-4-(3-oxo-2-aza-bicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide, ESI 537.

5 Example 7a

2-(3-N,N-Diethylaminocarbonylphenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, ESI 500, is prepared in accordance with the following scheme:

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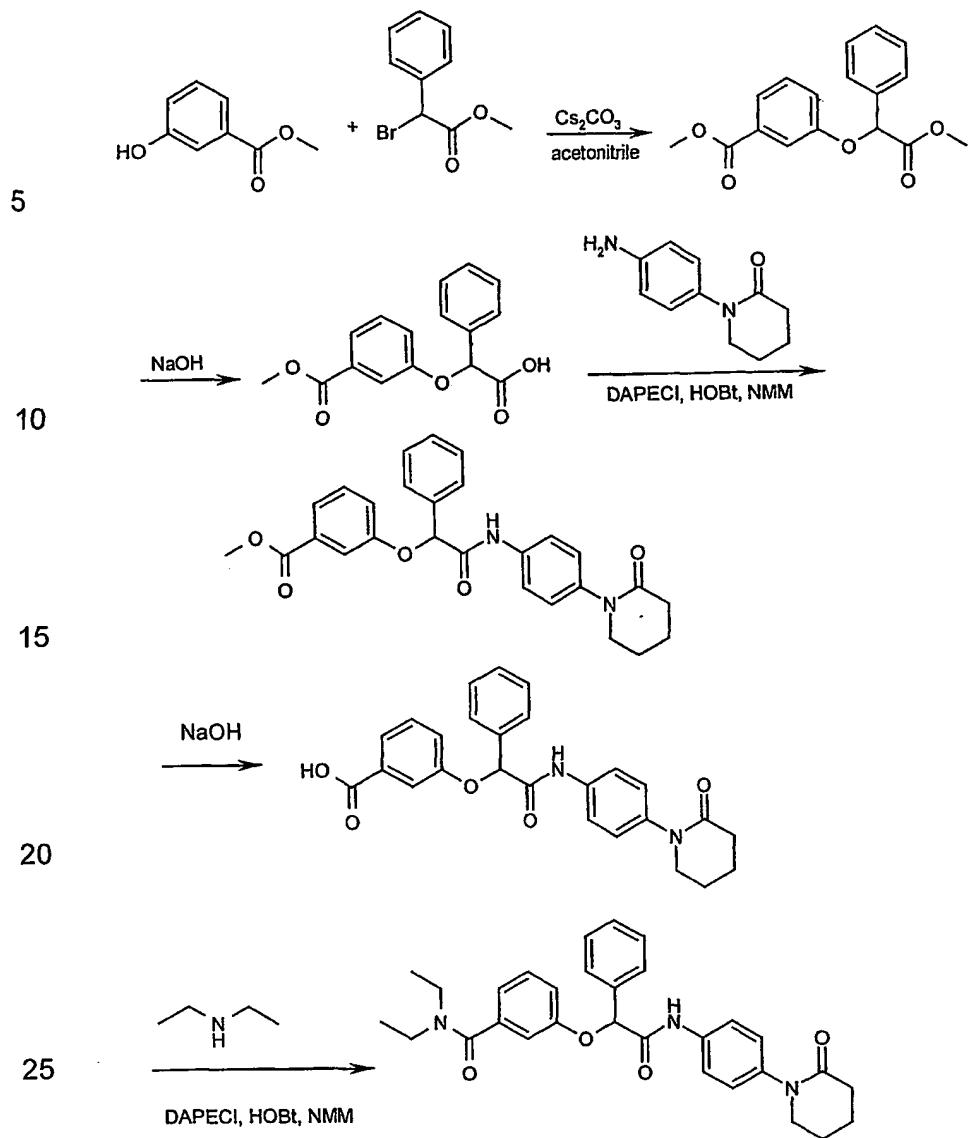
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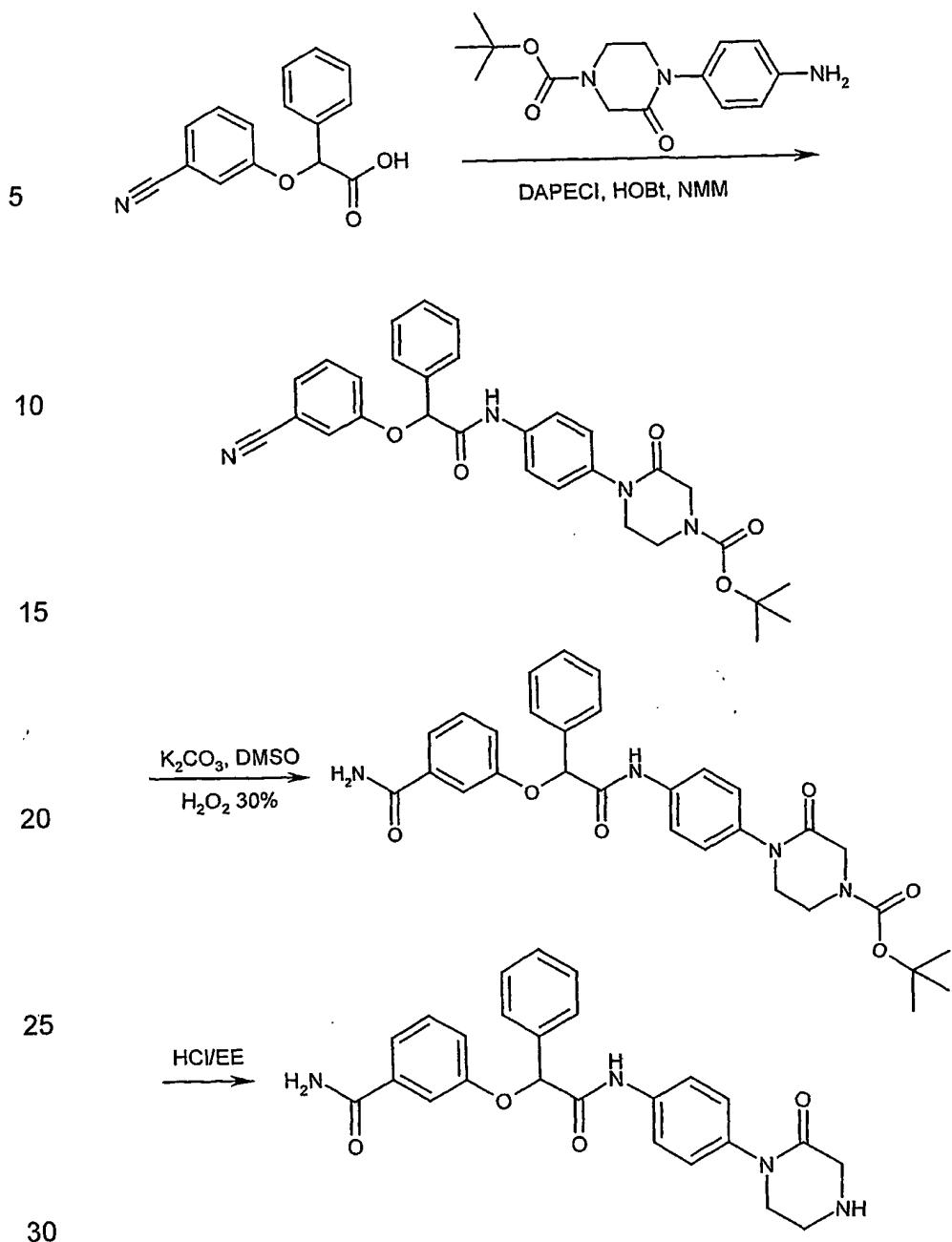
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Example 7b

2-(3-Aminocarbonylphenoxy)-N-[4-(2-oxopiperazin-1-yl)phenyl]-2-phenylacetamide, ESI 445, is prepared in accordance with the following scheme:



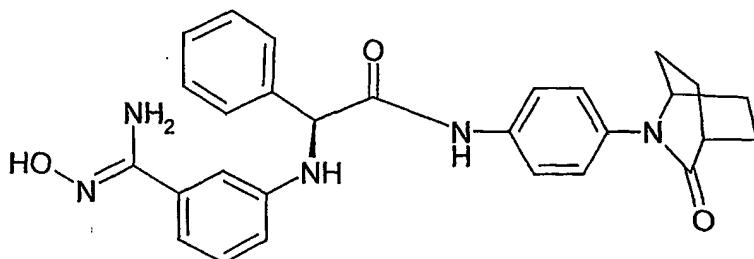
Example 8

The following compounds are obtained analogously to Example 2:

- 2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide, ESI 458;
- 2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, ESI 459;
- 5 2-[3-(*N*-hydroxyamidino)phenyl]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-propionamide, ESI 381;
- 10 2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]acetamide, ESI 382;
- 15 2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-valeramide, ESI 425;
- (2R)-[2-(3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-caprolactam-1-yl)phenyl]-2-phenylacetamide, ESI 472;
- 15 2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]-2-(2-trifluoromethylphenyl)acetamide, ESI 526;
- 20 2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide, ESI 490;
- 25 2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]-2-(2-thienyl)acetamide, ESI 464;
- 2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)-phenyl]valeramide, ESI 439;
- 30 (2S)-2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[3-fluoro-4-(2-oxo-piperidin-1-yl)phenyl]-2-phenylacetamide, ESI 476;
- 35 3-[3-(*N*-hydroxyamidino)phenyl]-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)-phenyl]propionamide, ESI 395;
- (2S)-2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]-4-methylvaleramide, ESI 438;
- 2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-valeramide, ESI 439;
- 35 2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxo-1*H*-pyrazin-1-yl)-phenyl]valeramide, ESI 422;

- (2S)-2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, hydrochloride, ESI 458;
- (2R)-2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide, ESI 439;
- 5 2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]valeramide, ESI 424;
- 10 2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide, ESI 452;
- 15 2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxopyridin-1-yl)phenyl]-valeramide, ESI 421;
- (2R)-2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-caprolactam-1-yl)phenyl]-4-methylvaleramide, ESI 452;
- 15 (2R)-2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]valeramide, ESI 425;
- 20 (2S)-2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]valeramide, ESI 425;
- 2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]-4-methylvaleramide, ESI 439;
- 2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]-3-methyl-4-trifluorobutyramide, ESI 478;
- 25 (2S)-2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)-benzyl]-2-phenylacetamide, ESI 472;
- (2S)-2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(3-oxo-2-aza-bicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide, ESI 484
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- 10 2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-butyramide, ESI 411;
- 15 2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-propionamide, ESI 397;
- 20 2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[3-trifluoromethyl-4-(3-oxo-2-azabicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide, ESI 552;
- 25 3-[3-(*N*-hydroxyamidino)phenyl]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylpropionamide, ESI 457;
- 30 3-[3-(*N*-hydroxyamidino)phenyl]-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylpropionamide, ESI 471;
- 35 2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[3-ethoxycarbonylmethoxy-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, ESI 561;
- 40 2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[3-ethoxycarbonylmethoxy-4-(2-oxopiperidin-1-yl)phenyl]valeramide, ESI 561;

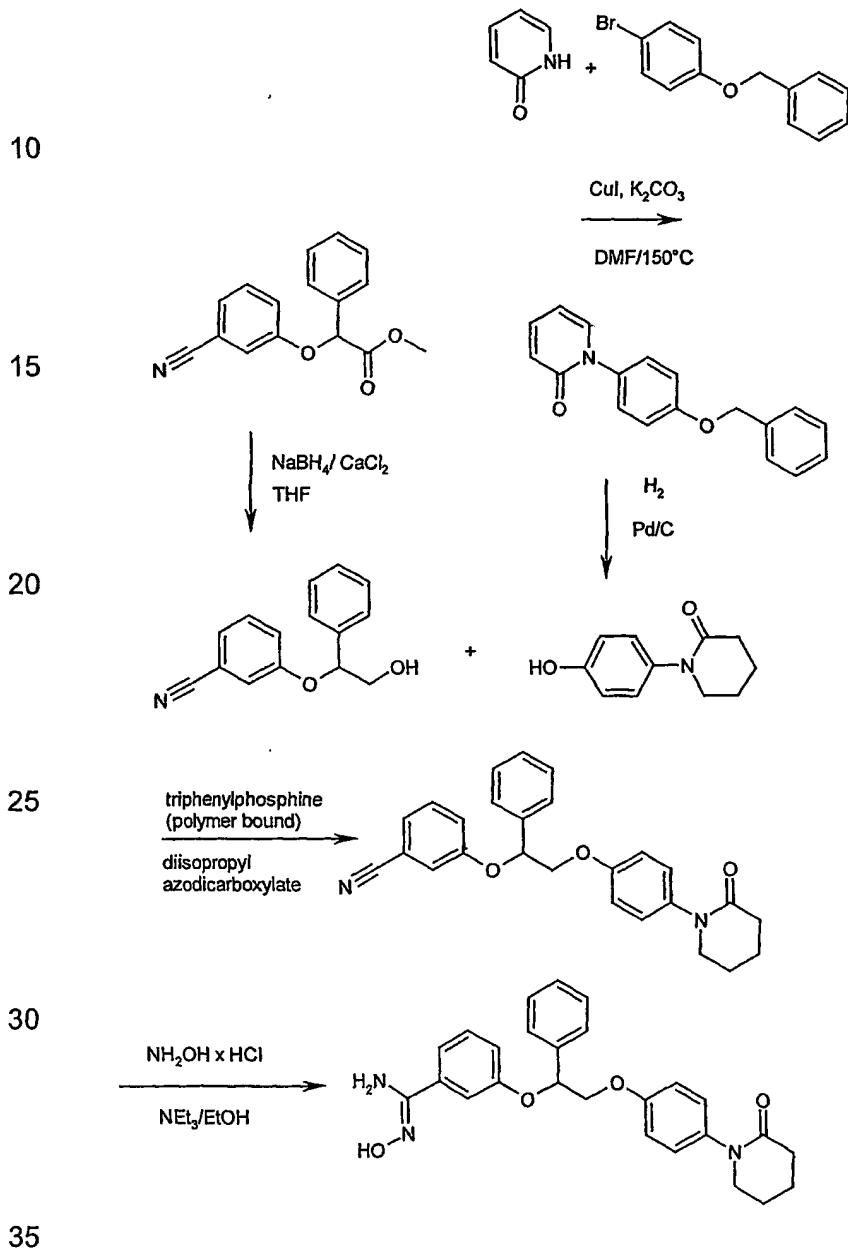
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Example 8a

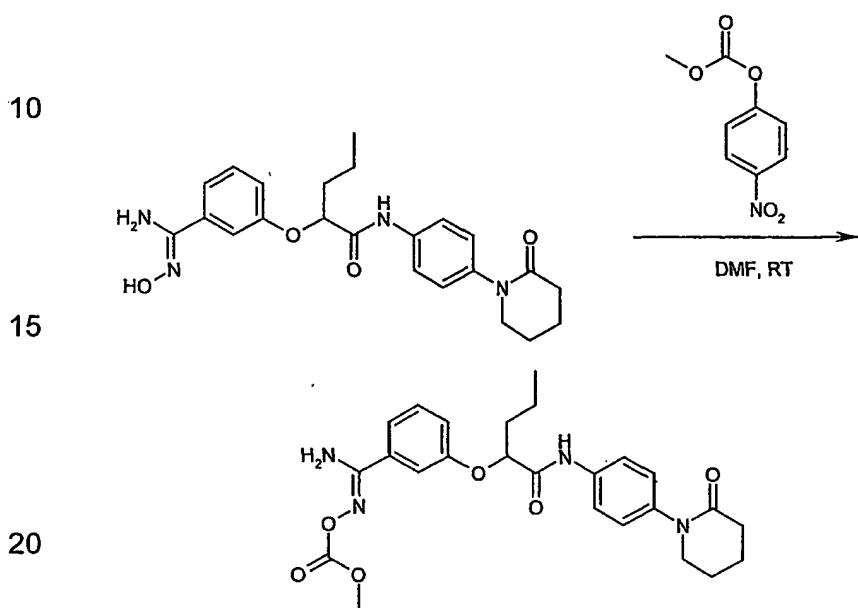
1-{4-[2-(3-(*N*-Hydroxyamidino)phenoxy)-2-phenylethoxy]phenyl}piperidin-2-one, ESI 446, is prepared in accordance with the following scheme:

5



Example 8b

5 2-[3-(*N*-ethoxycarbonyloxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)-
phenyl]valeramide, ESI 483, is prepared in accordance with the
following scheme:

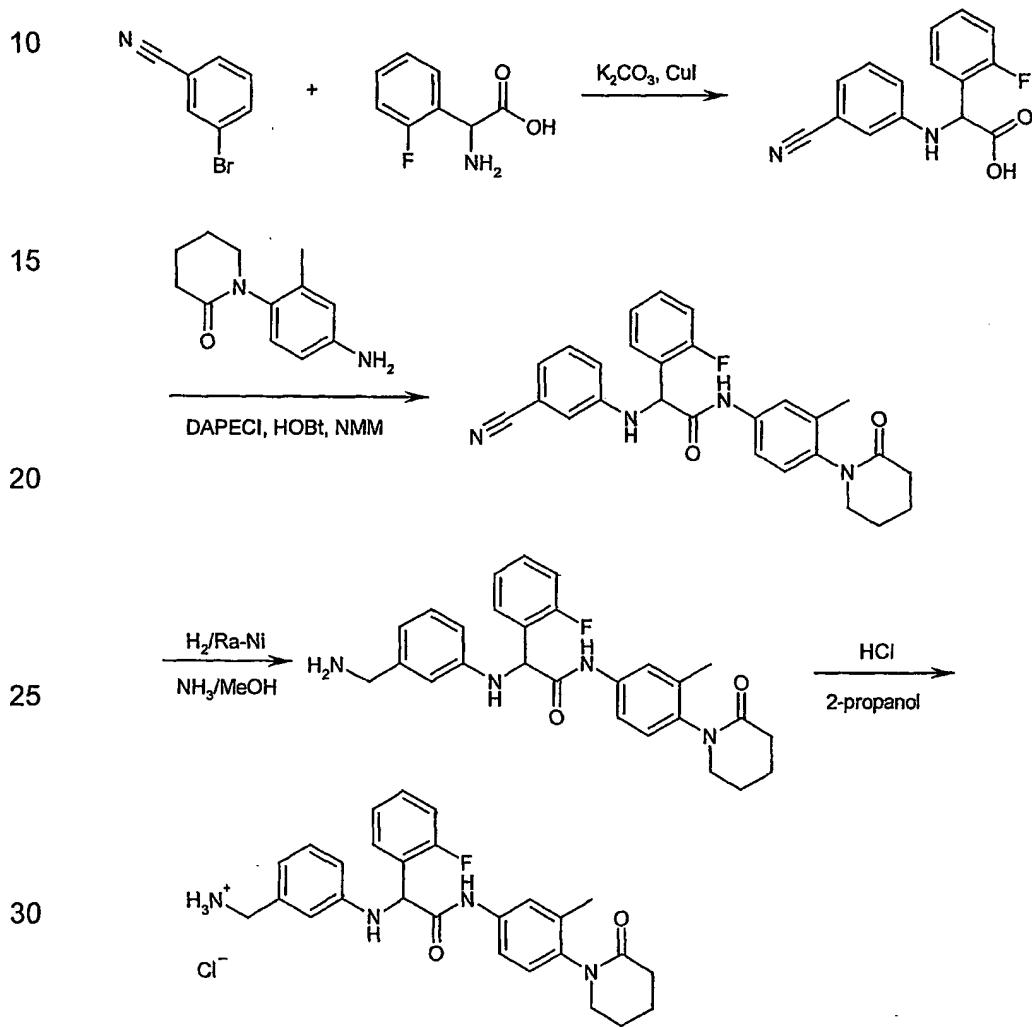


25 The following compounds are obtained analogously:

- 26 2-[3-(*N*-ethoxycarbonyloxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]valeramide, ESI 497;
- 30 (2*S*)-2-[3-(*N*-ethoxycarbonyloxyamidino)phenylamino]-*N*-[4-(2-oxo-piperidin-1-yl)phenyl]-4-methylvaleramide, ESI 494.

Example 9

5 2-(3-Aminomethylphenylamino)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-
 2-(2-fluorophenyl)acetamide is prepared as indicated in the following
 scheme:



35 1. 6.5 g (47.0 mmol) of potassium carbonate and 597 mg (3.10 mmol) of copper(I) iodide are added to a solution of 5.70 g (31.3 mmol)

of 3-bromobenzonitrile and 5.30 g (31.3 mmol) of 2-fluoro-dl-alpha-phenylglycine in 30 ml of N,N-dimethylacetamide, and the mixture is stirred at 90°C for 48 hours. The reaction mixture is then poured into 80 ml of 1N HCl and extracted twice with diethyl ether. The combined organic phases are extracted twice with 50 ml of 1N NaOH each time. The two basic aqueous phases are combined, acidified using 25% HCl and extracted twice with 80 ml of diethyl ether each time. These two organic phases are combined, dried using sodium sulfate and subsequently evaporated, giving (3-cyanophenylamino)-2-fluorophenylacetic acid as a yellowish solid; ESI 270.

2. 225 µl (2.22 mmol) of 4-methylmorpholine are added to a solution of 600 mg (2.22 mmol) of (3-cyanophenylamino)-2-fluorophenylacetic acid, 453 mg (2.22 mmol) of 1-(4-amino-2-methylphenyl)piperidin-2-one, 426 mg (2.22 mmol) of *N*-(3-dimethylaminopropyl)-*N'*-ethylcarbodiimide hydrochloride (DAPECI) and 300 mg (2.22 mmol) of hydroxybenzotriazole hydrate (HOBt) in 1 ml of DMF, and the mixture is stirred at room temperature for 24 hours. The reaction mixture is introduced into water, and the precipitate is filtered off, giving 2-(3-cyanophenylamino)-2-(2-fluorophenyl)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]acetamide as a colourless solid; ESI 456.

3. 300 mg of water-moist Raney nickel and 2.0 ml of methanolic ammonia solution are added to a solution of 660 mg (1.45 mmol) of 2-(3-cyanophenylamino)-2-(2-fluorophenyl)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]acetamide in 6 ml of methanol, and the mixture is hydrogenated at 60°C and 5 bar for 18 hours. The reaction mixture is filtered, and the filtrate is evaporated, giving 2-(3-cyanophenylamino)-2-(2-fluorophenyl)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]acetamide as a colourless solid; ESI 460.

4. 1.87 ml of 0.1N HCl in 2-propanol are added to 86 mg
(0.187 mmol) of 2-(3-cyanophenylamino)-2-(2-fluorophenyl)-N-[3-
5 methyl-4-(2-oxopiperidin-1-yl)phenyl]acetamide, and the mixture is left
to stand at room temperature for 2 hours. The reaction mixture is
evaporated and subsequently lyophilised, giving 2-(3-cyanophenyl-
10 amino)-2-(2-fluorophenyl)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-
acetamide, hydrochloride, as a colourless solid; ESI 460.

The following compounds are obtained analogously:

15 2-(3-aminomethylphenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-
phenylacetamide, hydrochloride, ESI 430;
20 2-(3-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide, ESI 447;
2-(3-aminomethylphenyl)-N-[4-(2-oxopiperidin-1-yl)phenyl]propion-
amide, ESI 352;
25 2-(3-aminomethylphenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]acet-
amide, ESI 353;
2-(3-aminomethylphenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]valer-
amide, ESI 396;
30 (2S)-2-(3-aminomethylphenylamino)-N-[4-(2-caprolactam-1-yl)-
phenyl]-2-phenylacetamide, ESI 443;
(2S)-2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-oxopiperidin-1-
yl)phenyl]acetamide, ESI 443;
35 2-(3-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-(2-trifluoromethylphenyl)acetamide, ESI 497;
(2S)-2-(3-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)-
phenyl]-2-phenylacetamide, ESI 429;

- (2R)-2-(3-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide, ESI 429;
- 2-(3-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-(2-methylphenyl)acetamide, ESI 443;
- 5 2-(3-aminomethylphenylamino)-N-[4-(2-caprolactam-1-yl)phenyl]-2-(2-fluorophenyl)acetamide, ESI 461;
- 10 2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-oxopiperidin-1-yl)-phenyl]-2-(2-methylphenyl)acetamide, ESI 457;
- 15 2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-oxopiperidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide, ESI 461;
- 20 2-(3-aminomethylphenoxy)-N-[3-methyl-4-(2-oxopiperidin-1-yl)-phenyl]valeramide, ESI 410;
- 25 2-(3-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-(2-thienyl)acetamide, ESI 435;
- 30 2-(3-aminomethylphenylamino)-N-[3-fluoro-4-(2-oxopiperidin-1-yl)-phenyl]-2-(3-fluorophenyl)acetamide, ESI 465;
- 35 (2S)-2-(3-aminomethylphenylamino)-N-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, ESI 447;
- (2S)-2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, hydrochloride, ESI 443;
- (2S)-2-(2-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide, ESI 429;
- 2-(2-aminomethylphenyl)-N-[4-(2-oxopiperidin-1-yl)phenyl]acetamide, ESI 338;
- 30 2-(2-aminomethylphenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, ESI 430;
- 2-(4-aminomethylphenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, ESI 430;
- 35 2-(3-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-(3-fluorophenyl)acetamide, ESI 447;

- 2-(3-aminomethylphenoxy)-N-[3-methyl-4-(2-caprolactam-1-yl)-phenyl]-2-phenylacetamide, ESI 458;
- (2R)-2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-caprolactam-1-yl)phenyl]-2-phenylacetamide, ESI 457;
- (2R)-2-(2-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide, ESI 429;
- (2S)-2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-caprolactam-1-yl)phenyl]-2-phenylacetamide, ESI 457;
- 2-(3-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-cyclohexylacetamide, ESI 435;
- 2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-oxopiperidin-1-yl)-phenyl]-2-cyclohexylacetamide, ESI 449;
- 2-(3-aminomethylphenylamino)-N-[4-(2-caprolactam-1-yl)phenyl]-4-methylvaleramide, ESI 423;
- (2R)-2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, hydrochloride, ESI 443;
- (2S)-2-(3-aminomethylphenylamino)-N-[4-(2-oxopyridin-1-yl)phenyl]-2-phenylacetamide, ESI 425;
- 2-(3-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-valeramide, ESI 395;
- 2-(3-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide, ESI 409;
- (2R)-2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide, ESI 423;
- (2R)-2-(3-aminomethylphenylamino)-N-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide, ESI 427;
- (2R)-2-(3-aminomethylphenylamino)-N-[4-(2-caprolactam-1-yl)-phenyl]-4-methylvaleramide, ESI 423;
- (2S)-2-(3-aminomethylphenylamino)-N-[2,5-dimethyl-4-(2-oxo-piperidin-1-yl)phenyl]-2-phenylacetamide, hydrochloride, ESI 457;

- 2-(3-aminomethylphenylamino)-N-[3-fluoro-4-(2-oxopiperidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 465;
- 2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-caprolactam-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 475;
- 5 2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-oxopiperidin-1-yl)-phenyl]-2-(2,4-difluorophenyl)acetamide, hydrochloride, ESI 479;
- 10 2-(3-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-(2,4-difluorophenyl)acetamide, hydrochloride, ESI 465;
- 10 2(S)-2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-oxopyrrolidin-1-yl)phenyl]-2-phenylacetamide, hydrochloride, ESI 429;
- 15 (2S)-2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide, ESI 423;
- 15 2-(3-aminomethylphenylamino)-N-methyl-N-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide, ESI 443;
- 20 2-(3-aminoethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, hydrochloride, ESI 443;
- 20 2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-oxopyrrolidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 447;
- 25 2-(3-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-4-trifluoro-3-methylbutyramide, ESI 449;
- 25 2-(3-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)benzyl]-2-phenylacetamide, ESI 443;
- (2S)-2-(3-aminomethylphenylamino)-N-[4-(3-oxo-2-azabicyclo[2.2.2]-oct-2-yl)phenyl]-2-phenylacetamide, ESI 455;
- 30 2-(3-aminomethylphenylamino)-N-[3-fluoro-4-(2-oxopyrrolidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 451;
- 30 2-(3-aminomethylphenylamino)-N-[4-(2-oxopyrrolidin-1-yl)phenyl]-2-phenylacetamide, hydrochloride, ESI 415;
- 35 2-(3-aminomethylphenylamino)-N-[4-(2-oxopyrrolidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 433;

- 2-(3-aminomethylphenylamino)-N-[4-ethyl-3-(2-oxopyrrolidin-1-yl)-phenyl]-2-phenylacetamide, ESI 443;
- (2S)-2-(3-aminomethylphenylamino)-N-[3-methoxy-4-(3-oxo-2-aza-bicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide, ESI 485;
- 5 2-(3-aminomethylphenylamino)-N-[3-fluoro-4-(2-caprolactam-1-yl)phenyl]-2-(2-fluorophenyl)acetamide, ESI 479;
- (2S)-2-(3-aminomethylphenylamino)-N-[3-fluoro-4-(2-caprolactam-1-yl)phenyl]-2-phenylacetamide, ESI 461;
- 10 (2S)-2-(3-aminomethylphenylamino)-N-[3-trifluoromethyl-4-(3-oxo-2-azabicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide, ESI 523;
- 2-(3-aminomethylphenylamino)-N-[3-methyl-4-(3-oxo-2-azabicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide, ESI 469;
- 15 (2S)-2-(3-aminomethylphenylamino)-N-[3-methoxy-4-(3-oxo-2-aza-bicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide, ESI 485;
- 2-(3-aminomethylphenylamino)-N-[3-trifluoromethyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, ESI 497;
- 20 2-(3-aminomethylphenylamino)-N-[3-methoxy-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, ESI 459;
- 2-(3-aminomethylphenylamino)-N-[3-methyl-4-(3-oxo-2-azabicyclo[2.2.2]oct-2-yl)phenyl]-2-(2-fluorophenyl)acetamide, ESI 487;
- 25 2-(3-aminomethyl-4-fluorophenylamino)-N-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide, ESI 447;
- 2-(3-aminomethylphenylamino)-N-[3-methyl-4-(3-oxo-2-azabicyclo[2.2.2]oct-2-yl)phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride,
- 30 ESI 487;
- 2-(3-aminomethylphenylamino)-N-[3-chloro-4-(2-oxopyrrolidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 467;
- 35 2-(3-aminomethylphenylamino)-N-[3-methoxy-4-(2-oxopiperidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide, ESI 477;

- 2-(3-aminomethylphenylamino)-N-[3-trifluoromethyl-4-(3-oxo-2-aza-bicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide, hydrochloride, ESI 523;
- 2-(3-aminomethylphenylamino)-N-[3-trifluoromethyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, hydrochloride, ESI 497;
- 5 3-(3-aminomethylphenyl)-N-[4-(2-caprolactam-1-yl)phenyl]-2-phenylpropionamide, hydrochloride, ESI 456;
- 2-(3-aminomethylphenoxy)-N-[3-methyl-4-(2-oxopyrrolidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 448;
- 10 3-(3-aminomethylphenyl)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylpropionamide, hydrochloride, ESI 428;
- 2-(3-aminomethylphenoxy)-N-[3-methyl-4-(2-caprolactam-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 476;
- 15 3-(3-aminomethylphenyl)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylpropionamide, hydrochloride, ESI 442;
- 2-(3-aminomethylphenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 448;
- 20 2-(3-aminomethylphenoxy)-N-[3-methyl-4-(2-oxopiperidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 462;
- 3-(3-aminomethylphenyl)-N-[3-methyl-4-(2-oxopyrrolidin-1-yl)phenyl]-2-phenylpropionamide, hydrochloride, ESI 428;
- 25 (2S)-2-(3-aminomethylphenylamino)-N-[4-(5,5-dimethyl-2-oxopyrrolidin-1-yl)phenyl]-2-phenylacetamide, hydrochloride, ESI 443;
- (2S)-2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-oxo-1,3-oxazinan-3-yl)phenyl]-2-phenylacetamide, hydrochloride, ESI 445;
- 30 (2S)-2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-oxooxazolidin-3-yl)phenyl]-2-phenylacetamide, hydrochloride, ESI 431;
- 3-(3-aminomethylphenyl)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)propionamide, hydrochloride, ESI 460;
- 35 3-(3-aminomethylphenyl)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)propionamide, hydrochloride, ESI 446;

- 3-(3-aminomethylphenylamino)-*N*-[3-chloro-4-(2-oxopiperidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 481;
- 3-(3-aminomethylphenylamino)-*N*-[3-chloro-4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide, hydrochloride, ESI 463;
- 5 3-(3-aminomethylphenoxy)-*N*-[3-chloro-4-(2-oxopiperidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 482;
- 10 3-(3-aminomethylphenyl)-*N*-[3-methyl-4-(2-oxopyrrolidin-1-yl)phenyl]-2-(2-fluorophenyl)propionamide, hydrochloride, ESI 446;
- 15 3-(3-aminomethylphenyl)-*N*-[3-chloro-4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)propionamide, hydrochloride, ESI 480;
- 20 3-(3-aminomethylphenylamino)-*N*-[4-(2,6-dioxopiperidin-1-yl)phenyl]-2-phenylacetamide, trifluoroacetate, ESI 443;
- 25 3-(3-aminomethylphenylamino)-*N*-[3-chloro-4-(2,5-dioxopyrrolidin-1-yl)phenyl]-2-phenylacetamide, ditrifluoroacetate, ESI 463;
- 30 2-(3-aminomethyl-4-fluorophenylamino)-*N*-[3-methyl-4-(3-oxo-2-aza-bicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide, hydrochloride, ESI 487;
- 35 2-[3-(N-tert-butoxycarbonylaminomethyl)phenylamino]-*N*-[3-chloro-4-(2,5-dioxopyrrolidin-1-yl)phenyl]-2-phenylacetamide, ESI 563;
- 2-(3-aminomethylphenylamino)-*N*-[3-methyl-4-(3-oxo-2-azabicyclo-[2.2.2]oct-2-yl)phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 487;
- 2-(3-aminomethylphenylamino)-*N*-[3-methyl-4-(2-oxooxazolidin-3-yl)-phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 449;
- 3-(3-aminomethylphenylamino)-*N*-[3-methoxy-4-(2-oxopiperidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 477;
- 3-(3-aminomethylphenylamino)-*N*-[3-chloro-4-(2-oxopyrrolidin-1-yl)-phenyl]-2-phenylacetamide, hydrochloride, ESI 449;
- 2-(3-aminomethylphenoxy)-*N*-[3-methyl-4-(3-oxo-2-azabicyclo[2.2.2]-oct-2-yl)phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 488;

3-(3-aminomethylphenylamino)-*N*-[4-(5,5-dimethyl-2-oxopyrrolidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 461;

5

3-(3-aminomethylphenoxy)-*N*-[4-(5,5-dimethyl-2-oxopyrrolidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 462;

10

2-(3-aminomethylphenoxy)-*N*-[3-methyl-4-(2-oxo-oxazolidin-3-yl)-phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 450;

15

3-(3-aminomethylphenoxy)-*N*-[3-methoxy-4-(2-oxopiperidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide, hydrochloride, ESI 478.

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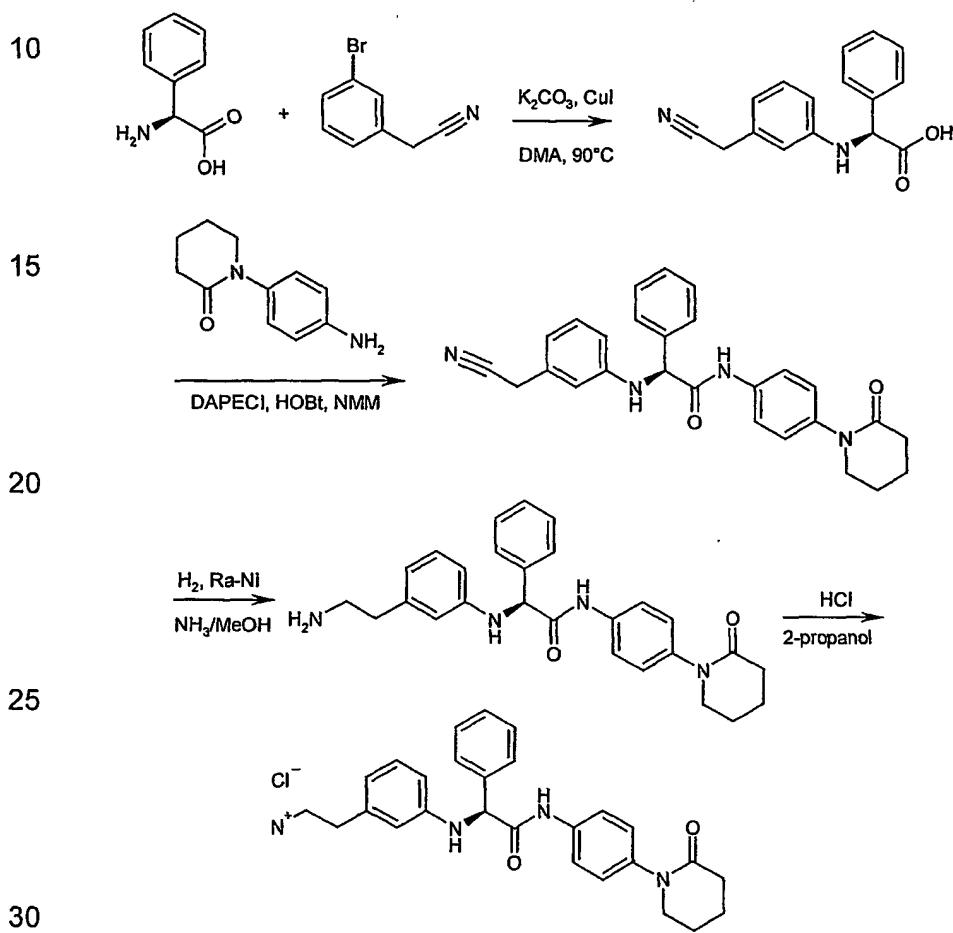
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Example 9a

5 (2S)-2-(3-Aminoethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, hydrochloride, ESI 443, is prepared in accordance with the following scheme:



Example 10

5 The following is obtained analogously to Example 3:

2-(3-amidinophenoxy)-N-[4-(2-oxopiperazin-1-yl)phenyl]-2-phenylacetamide, diformate, ESI 444.

10 Example 10a

15 2-(3-Amidinophenoxy)-N-[3-carboxymethoxy-4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide is prepared as indicated in the following scheme:

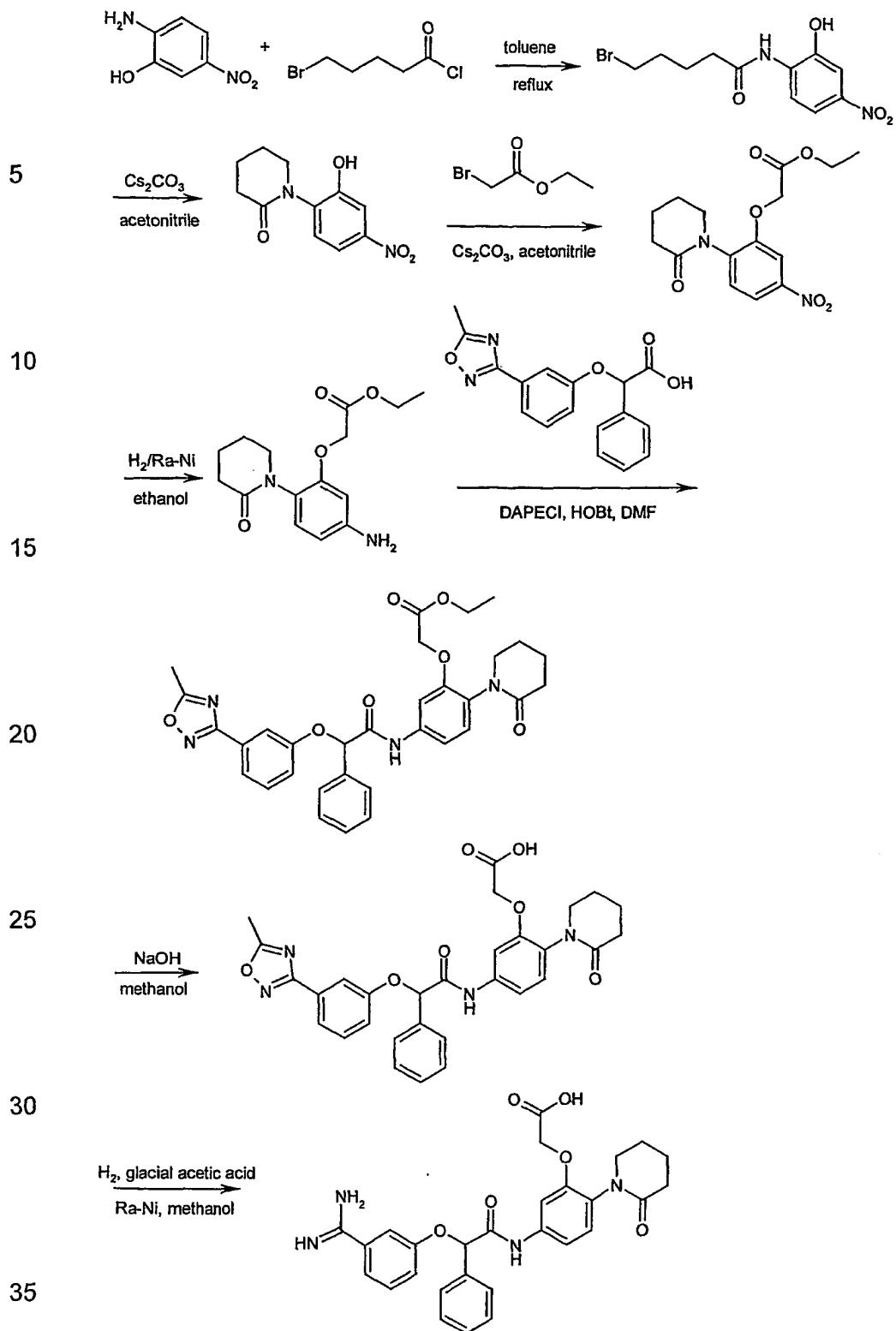
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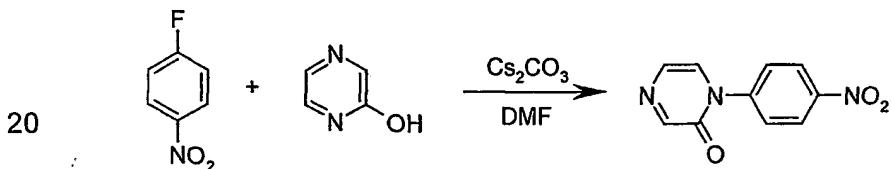
The following compounds are obtained analogously:

- 5 2-(3-amidinophenoxy)-N-[3-ethoxycarbonylmethoxy-4-(2-oxo-piperidin-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 545;
- 10 2-(3-amidinophenoxy)-N-[3-ethoxycarbonylmethoxy-4-(2-oxo-piperidin-1-yl)phenyl]valeramide, acetate, ESI 511;
- 15 3-(3-amidinophenoxy)-N-[3-carboxymethoxy-4-(2-oxopiperidin-1-yl)phenyl]valeramide, acetate, ESI 483;
- 20 3-(3-amidinophenoxy)-N-[3-carboxymethoxy-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, acetate, ESI 517.

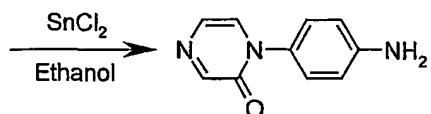
11. Examples of the preparation of intermediates

15

11.1 1-(4-Aminophenyl)-1*H*-pyrazin-2-one



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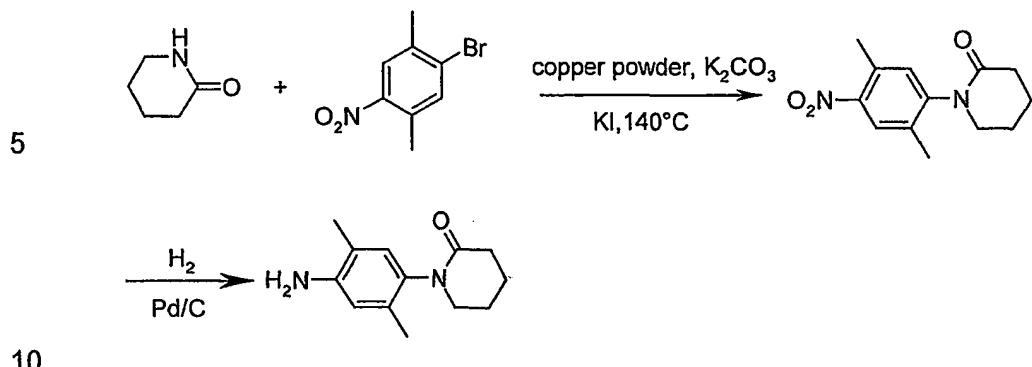


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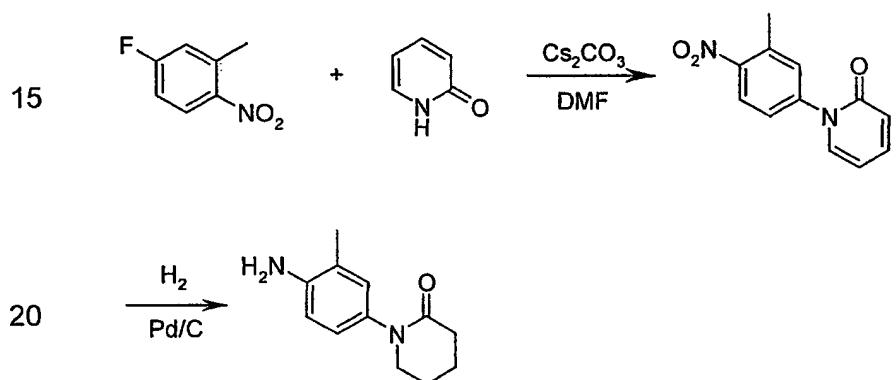
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- 82 -

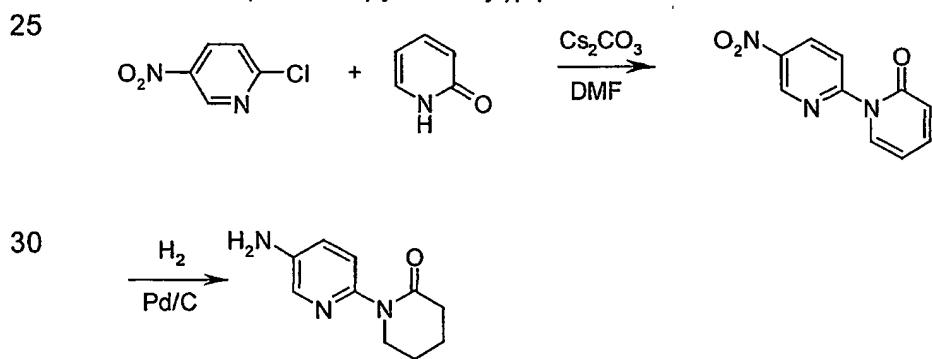
11.2 1-(4-Amino-2,5-dimethylphenyl)piperidin-2-one



11.3 1-(4-Amino-3-methylphenyl)piperidin-2-one

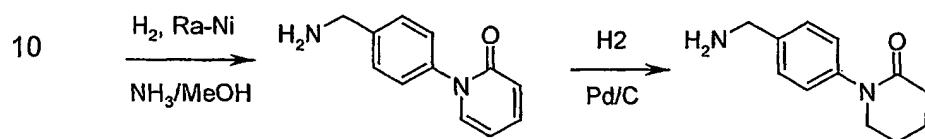
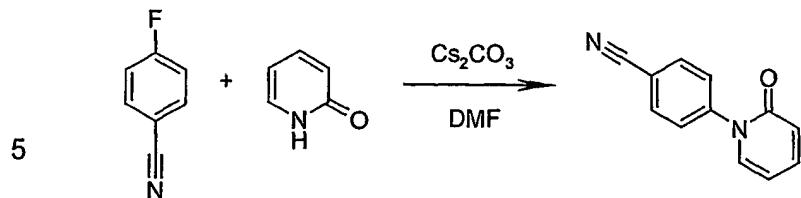


11.4 1-(5-Aminopyridin-2-yl)piperidin-2-one

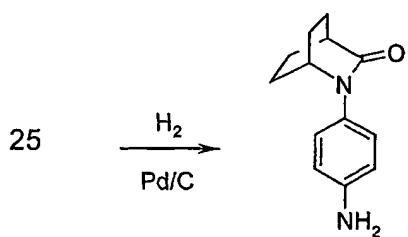
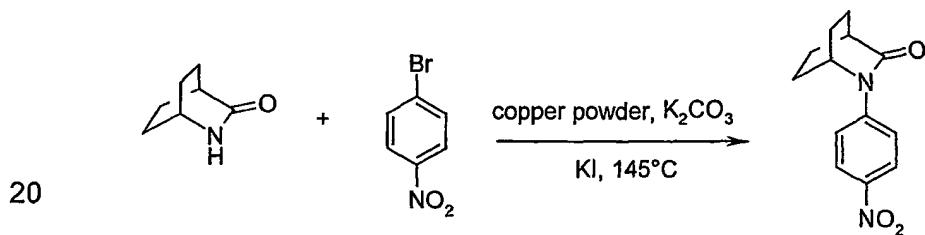


- 83 -

11.5 1-(4-Aminomethylphenyl)piperidin-2-one



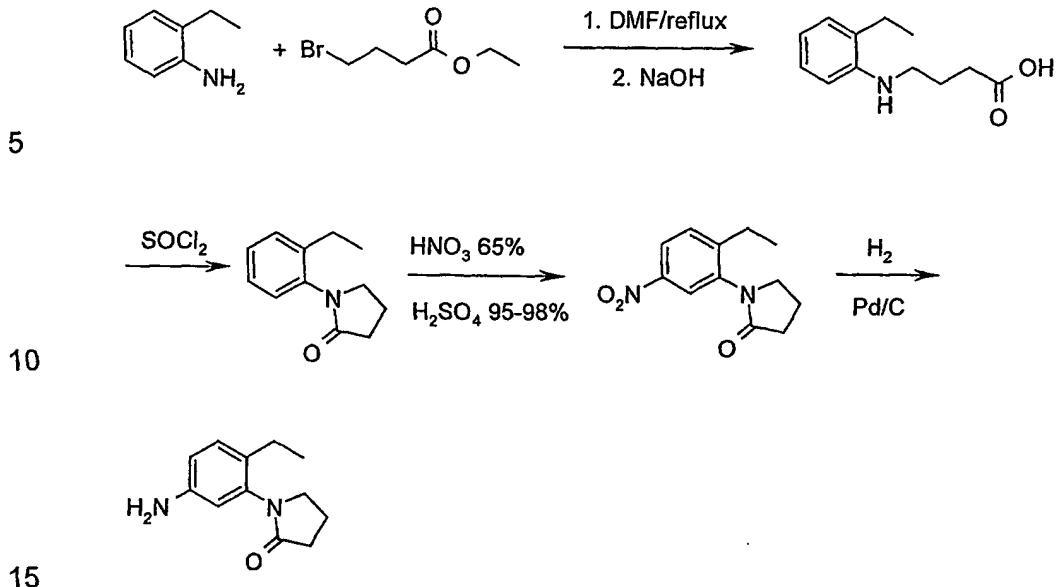
11.6 2-(4-Aminophenyl)-2-azabicyclo[2.2.2]octan-3-one



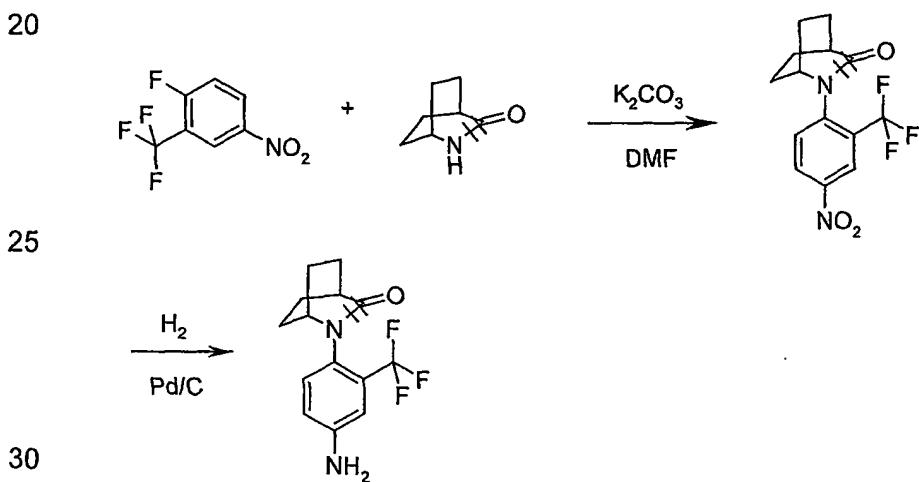
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11.7 1-(3-Amino-6-ethylphenyl)pyrrolidin-2-one

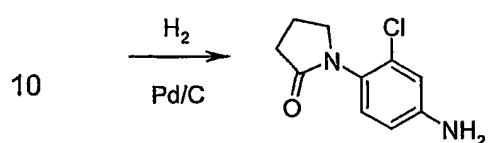
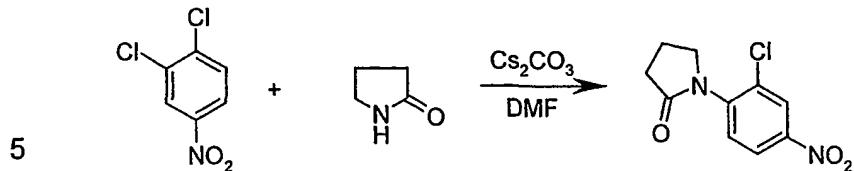


11.8 2-(4-Amino-2-trifluoromethylphenyl)-2-azabicyclo[2.2.2]octan-3-one



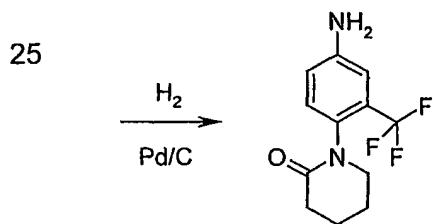
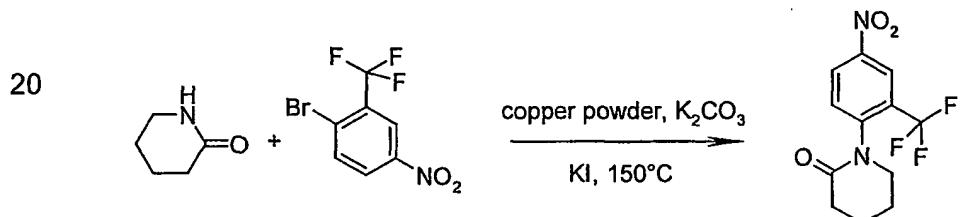
- 85 -

11.9 1-(4-Amino-3-chlorophenyl)pyrrolidin-2-one



15

11.11 1-(4-Amino-2-trifluoromethylphenyl)piperidin-2-one

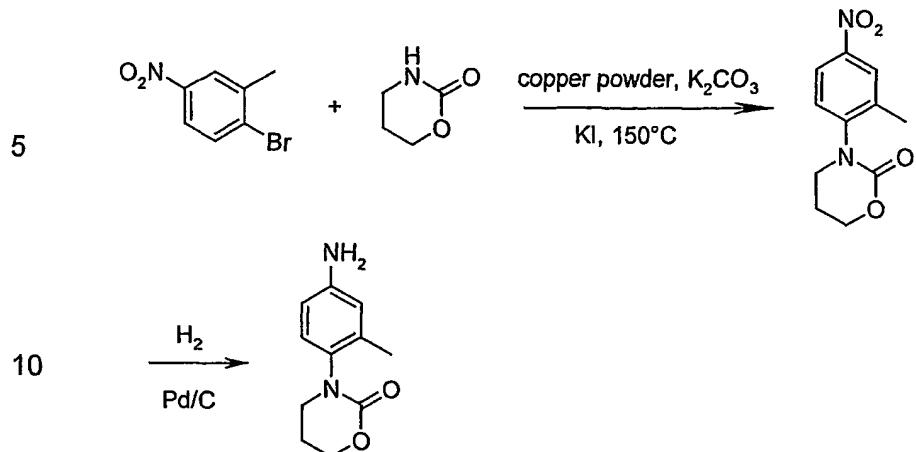


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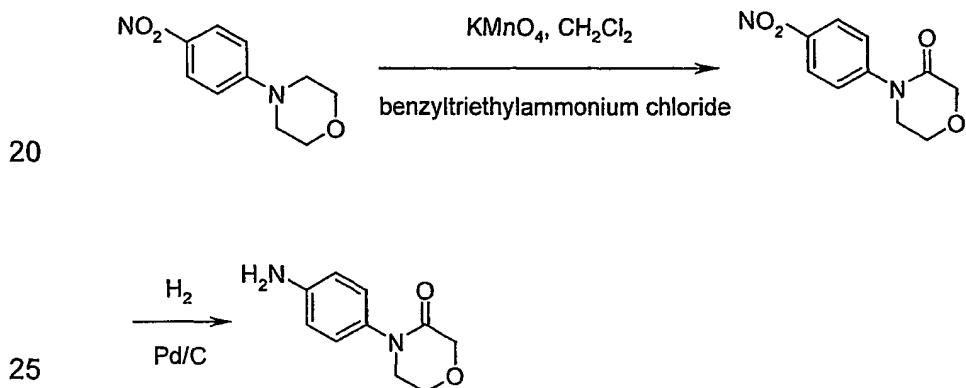
- 86 -

11.12 3-(4-Amino-2-methylphenyl)-1,3-oxazinan-2-one



15

11.13 4-(4-Aminophenyl)morpholin-3-one

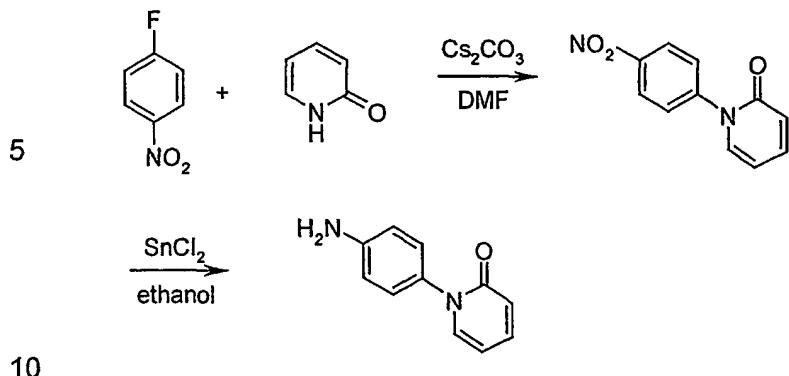


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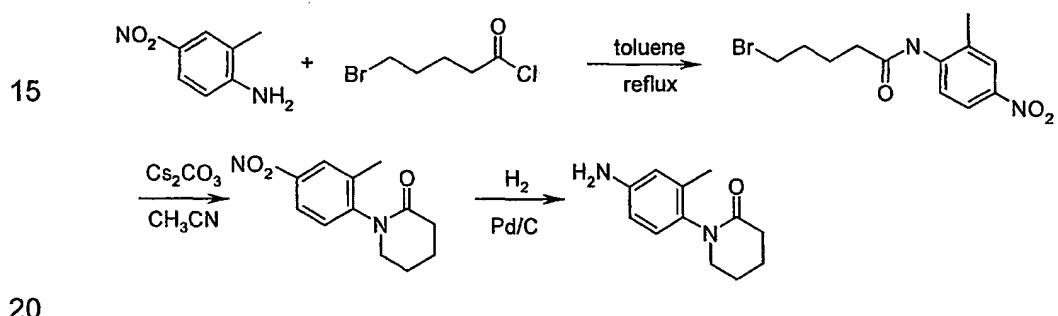
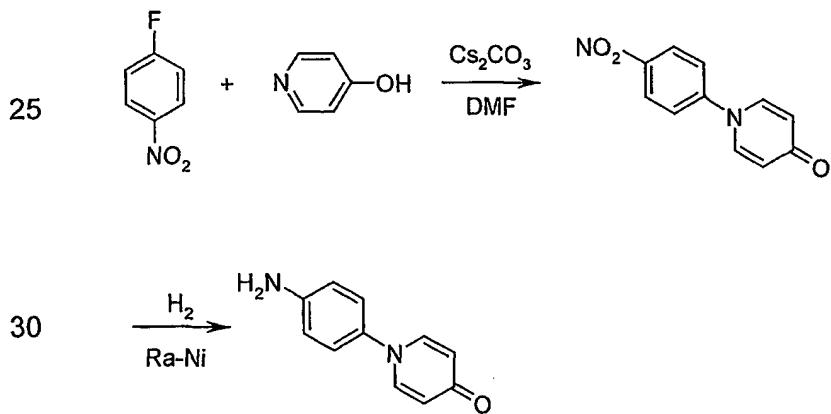
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- 87 -

11.14 1-(4-Aminophenyl)pyridin-2-one

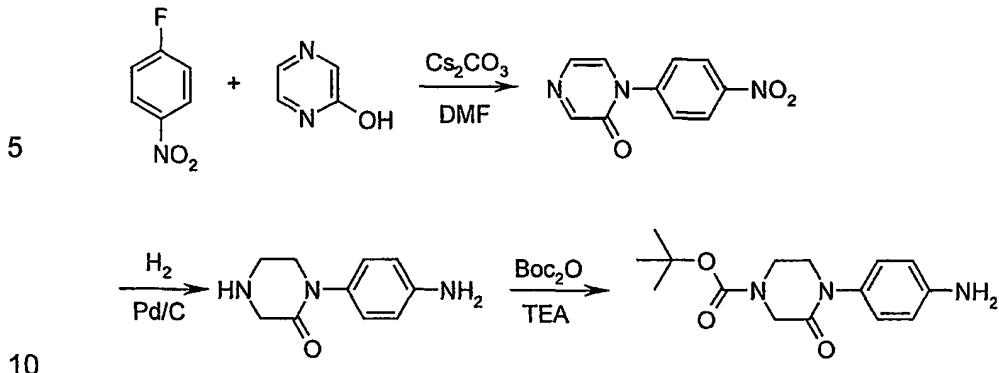


11.15 1-(4-Amino-2-methylphenyl)piperidin-2-one

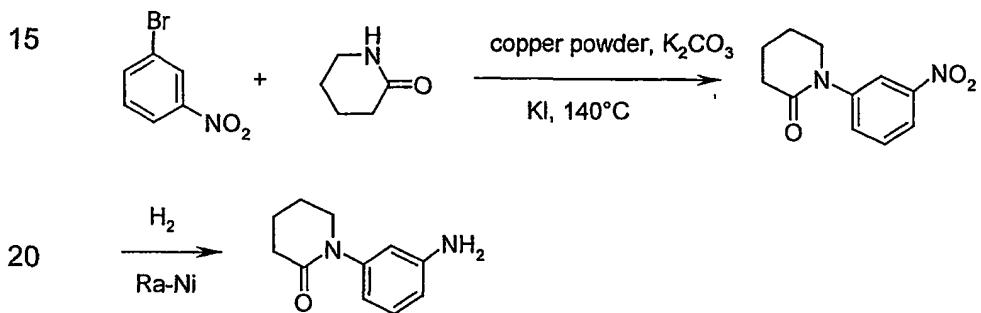
11.16 1-(4-Aminophenyl)-1*H*-pyridin-4-one

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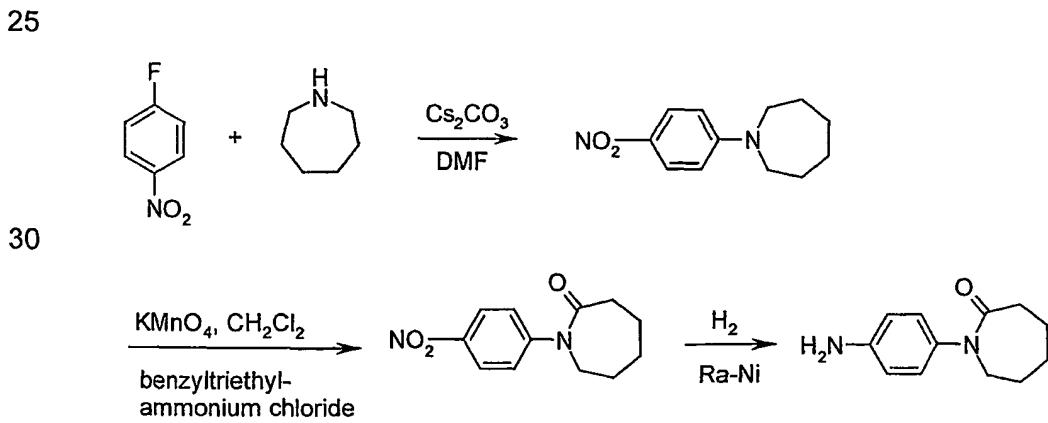
11.17 1-(4-Aminophenyl)-4-tert-butoxycarbonylpiperazin-2-one



11.18 1-(3-Aminophenyl)piperidin-2-one

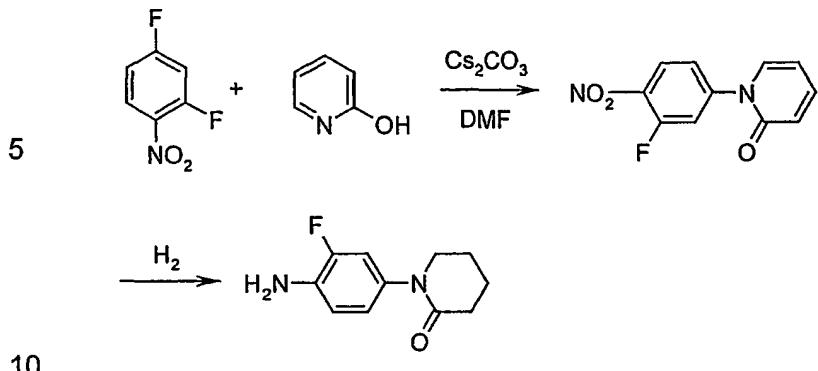


11.19 1-(4-Aminophenyl)-2-caprolactam

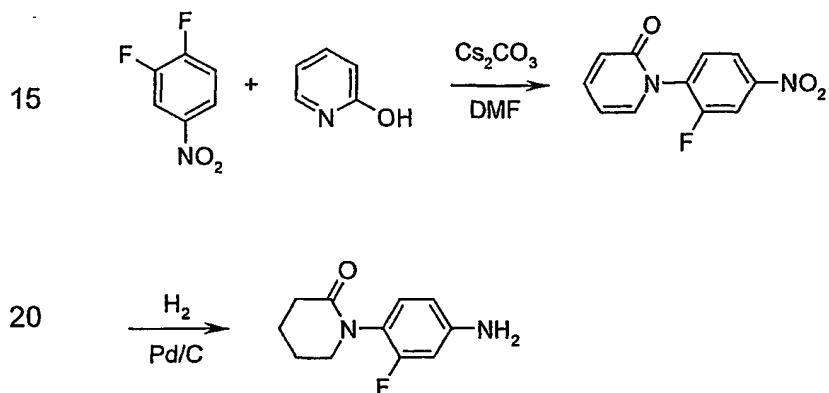


- 89 -

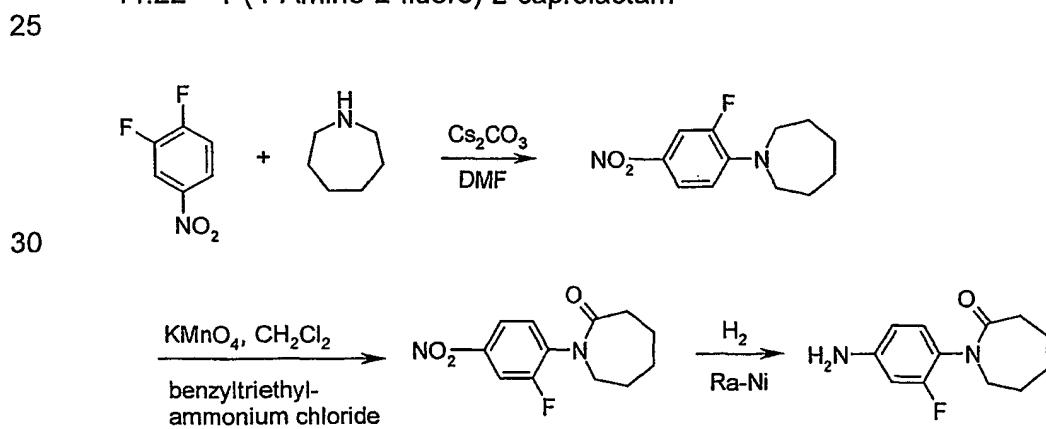
11.20 1-(4-Amino-3-fluorophenyl)piperidin-2-one



11.21 1-(4-Amino-2-fluorophenyl)piperidin-2-one

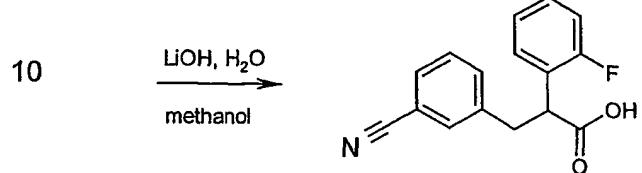
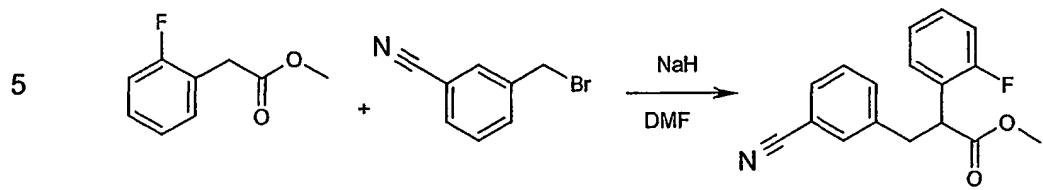


11.22 1-(4-Amino-2-fluoro)-2-caprolactam



- 90 -

11.23 2-(2-Fluorophenyl)-3-(3-cyanophenyl)propionic acid



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The examples below relate to pharmaceutical preparations:

Example A: Injection vials

- 5 A solution of 100 g of an active ingredient of the formula I and 5 g of disodium hydrogenphosphate in 3 l of bidistilled water is adjusted to pH 6.5 using 2N hydrochloric acid, sterile filtered, transferred into injection vials, lyophilised under sterile conditions and sealed under sterile conditions. Each injection vial contains 5 mg of active ingredient.

10

Example B: Suppositories

- A mixture of 20 g of an active ingredient of the formula I is melted with 100 g of soya lecithin and 1400 g of cocoa butter, poured into moulds and allowed to cool. Each suppository contains 20 mg of active ingredient.

15

Example C: Solution

- 20 A solution is prepared from 1 g of an active ingredient of the formula I, 9.38 g of $\text{NaH}_2\text{PO}_4 \cdot 2 \text{ H}_2\text{O}$, 28.48 g of $\text{Na}_2\text{HPO}_4 \cdot 12 \text{ H}_2\text{O}$ and 0.1 g of benzalkonium chloride in 940 ml of bidistilled water. The pH is adjusted to 6.8, and the solution is made up to 1 l and sterilised by irradiation. This solution can be used in the form of eye drops.

25

Example D: Ointment

- 500 mg of an active ingredient of the formula I are mixed with 99.5 g of Vaseline under aseptic conditions.

30

Example E: Tablets

- 35 A mixture of 1 kg of active ingredient of the formula I, 4 kg of lactose, 1.2 kg of potato starch, 0.2 kg of talc and 0.1 kg of magnesium stearate is pressed in a conventional manner to give tablets in such a way that each tablet contains 10 mg of active ingredient.

Example F: Coated tablets

Tablets are pressed analogously to Example E and subsequently coated in
5 a conventional manner with a coating of sucrose, potato starch, talc, tragacanth and dye.

Example G: Capsules

10 2 kg of active ingredient of the formula I are introduced in a conventional manner into hard gelatine capsules in such a way that each capsule contains 20 mg of the active ingredient.

Example H: Ampoules

15 A solution of 1 kg of active ingredient of the formula I in 60 l of bidistilled water is sterile filtered, transferred into ampoules, lyophilised under sterile conditions and sealed under sterile conditions. Each ampoule contains 10 mg of active ingredient.

20

25

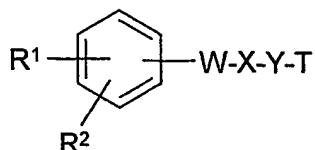
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Patent Claims

1. Compounds of the formula I

5



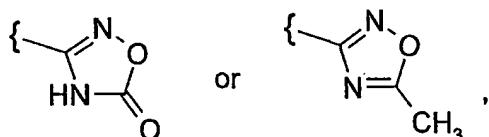
10

in which

15

R^1 is CN , $CON(R^3)_2$, $-[C(R^4)_2]_nN(R^3)_2$,
 $-C(=NH)-NH_2$ which is unsubstituted or monosubstituted by
 $C(=O)R^3$, $COOR^3$, OR^3 , $OCOR^3$, $OCOOR^3$ or by a conven-
tional amino-protecting group, or is

20



30

R^2 is H , Hal , A , OR^3 , $N(R^3)_2$, NO_2 , CN , $COOR^3$, $CON(R^3)_2$,
 $-[C(R^4)_2]_n-Ar$, $-[C(R^4)_2]_n-Het$ or $-[C(R^4)_2]_n-cycloalkyl$,

R^3 is H , A , $-[C(R^4)_2]_n-Ar$, $-[C(R^4)_2]_n-Het$ or $-[C(R^4)_2]_n-cycloalkyl$,

R^4 is H or A ,

W is $-C(R^3)_2-$, $-[C(R^3)_2]_2-$, $-OC(R^3)_2-$ or $-NR^3C(R^3)_2-$,

X is $CONR^3$, $CONR^3C(R^4)_2-$, $-C(R^4)_2NR^3-$, $-C(R^4)_2NR^3C(R^4)_2-$,
 $-C(R^4)_2O-$ or $-C(R^4)_2OC(R^4)_2-$,

Y is alkylene, cycloalkylene, Het-diyl or Ar-diyl,

T is OR^3 , $N(R^3)_2$ or a monocyclic or bicyclic, saturated, unsatu-
rated or aromatic heterocyclic radical having from 1 to 4 N, O
and/or S atoms which is unsubstituted or monosubstituted,
disubstituted or trisubstituted by Hal , A , $-[C(R^4)_2]_n-Ar$,

35

-

-[C(R⁴)₂]_n-Het, -[C(R⁴)₂]_n-cycloalkyl, OR³, N(R³)₂, NO₂, CN, COOR³, CON(R³)₂, NR³COA, NR³SO₂A, COR³, SO₂NR³, S(O)_mA and/or carbonyl oxygen,

5 A is unbranched or branched alkyl having 1-6 carbon atoms, in which one or two CH₂ groups may be replaced by O or S atoms and/or by -CH=CH- groups and/or, in addition, 1-7 H atoms may be replaced by F,

10 Ar is phenyl, naphthyl or biphenyl, each of which is unsubstituted or monosubstituted, disubstituted or trisubstituted by Hal, A, OR⁴, N(R⁴)₂, NO₂, CN, COOR⁴, CON(R⁴)₂, NR⁴COA, NR⁴SO₂A, COR⁴, SO₂NR⁴, S(O)_mA, -[C(R⁴)₂]_n-COOR³ or O-[C(R⁴)₂]_o-COOR³,

15 Het is a monocyclic or bicyclic, saturated, unsaturated or aromatic heterocyclic radical having from 1 to 4 N, O and/or S atoms which is unsubstituted or monosubstituted, disubstituted or trisubstituted by Hal, A, -[C(R⁴)₂]_n-Ar, -[C(R⁴)₂]_n-Het', -[C(R⁴)₂]_n-cycloalkyl, OR³, N(R³)₂, NO₂, CN, COOR³, CON(R³)₂, NR³COA, NR³SO₂A, COR³, SO₂NR³, S(O)_mA and/or carbonyl oxygen,

20 Het' is a monocyclic or bicyclic, saturated, unsaturated or aromatic heterocyclic radical having from 1 to 4 N, O and/or S atoms which is unsubstituted or monosubstituted, disubstituted or trisubstituted by Hal, A, OR³, N(R³)₂, NO₂, CN, COOR³, CON(R³)₂, NR³COA, NR³SO₂A, COR³, SO₂NR³, S(O)_mA and/or carbonyl oxygen,

25 Hal is F, Cl, Br or I,

30 m and are each, independently of one another, 0, 1 or 2,

 n are each, independently of one another, 0, 1 or 2,

 o is 1, 2 or 3,

35 and their pharmaceutically usable derivatives, solvates and stereoisomers, including mixtures thereof in all ratios.

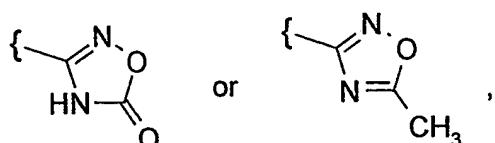
2. Compounds according to Claim 1, in which

R² is H,

5 and their pharmaceutically usable derivatives, solvates and stereo-isomers, including mixtures thereof in all ratios.

3. Compounds according to Claim 1, in which

10 R¹ is -C(=NH)-NH₂ which is unsubstituted or monosubstituted by OH, or is



15 and their pharmaceutically usable derivatives, solvates and stereo-isomers, including mixtures thereof in all ratios.

4. Compounds according to Claim 1, in which

20 Ar is phenyl which is unsubstituted or monosubstituted or disubstituted by Hal, A, OR⁴ or O-[C(R⁴)₂]-COOR³, and their pharmaceutically usable derivatives, solvates and stereo-isomers, including mixtures thereof in all ratios.

25

5. Compounds according to Claim 1, in which

Het is a monocyclic or bicyclic, saturated, unsaturated or aromatic heterocyclic radical having 1 to 2 N and/or O atoms which is monosubstituted or disubstituted by A and/or carbonyl oxygen,
30 and their pharmaceutically usable derivatives, solvates and stereo-isomers, including mixtures thereof in all ratios.

35

6. Compounds according to Claim 1, in which

W is $-\text{OC}(\text{R}^3)_2-$ or $-\text{NR}^3\text{C}(\text{R}^3)_2-$,
and their pharmaceutically usable derivatives, solvates and stereo-
isomers, including mixtures thereof in all ratios.

5

7. Compounds according to Claim 1, in which

W is $-\text{OC}(\text{R}^3)_2-$ or $-\text{NR}^3\text{C}(\text{R}^3)_2-$,

R³ is H, A or $-(\text{CH}_2)_n\text{-Ar}$,

10

Ar is phenyl which is unsubstituted or monosubstituted or
disubstituted by Hal, A, OR⁴ or O-[C(R⁴)₂]-COOR³,

n is 0 or 1,

and their pharmaceutically usable derivatives, solvates and stereo-
isomers, including mixtures thereof in all ratios.

15

8. Compounds according to Claim 1, in which

W is $-\text{OCHR}^3-$ or $-\text{NHCHR}^3-$,

R³ is H, A or $-(\text{CH}_2)_n\text{-Ar}$,

20

Ar is phenyl which is unsubstituted or monosubstituted or
disubstituted by Hal, A, OR⁴ or O-[C(R⁴)₂]-COOR³,

n is 0 or 1,

25

and their pharmaceutically usable derivatives, solvates and stereo-
isomers, including mixtures thereof in all ratios.

9. Compounds according to Claim 1, in which

X is CONH or CONH(CH₂)₂-,

30

and their pharmaceutically usable derivatives, solvates and stereo-
isomers, including mixtures thereof in all ratios.

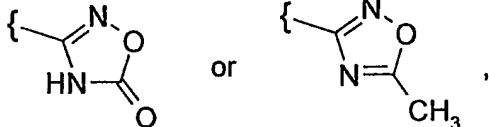
10. Compounds according to Claim 1, in which

35

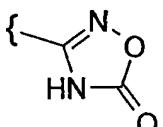
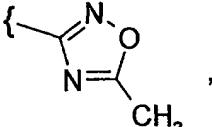
Y is alkylene or Ar-diyl,

Ar is phenyl which is unsubstituted or monosubstituted or disubstituted by Hal, A, OR⁴ or O-[C(R⁴)₂]-COOR³, and their pharmaceutically usable derivatives, solvates and stereoisomers, including mixtures thereof in all ratios.

5

11. Compounds according to Claim 1, in which
- T is N(R³)₂ or a monocyclic or bicyclic, saturated, unsaturated or aromatic heterocyclic radical having 1 to 2 N and/or O atoms which is monosubstituted or disubstituted by A and/or carbonyl oxygen,
- R³ is H or A,
- and their pharmaceutically usable derivatives, solvates and stereoisomers, including mixtures thereof in all ratios.
- 15
12. Compounds according to Claim 1, in which
- R¹ is -C(=NH)-NH₂ which is unsubstituted or monosubstituted by OH, or is
- 20
- 
- R² is H,
- R³ is H, A or -(CH₂)_n-Ar,
- R⁴ is H or A,
- W is -OC(R³)₂- or -NR³C(R³)₂-,
- X is CONH or CONH(CH₂)₂,
- Y is alkylene or Ar-diyl,
- 30
- T is N(R³)₂ or a monocyclic or bicyclic, saturated, unsaturated or aromatic heterocyclic radical having 1 to 2 N and/or O atoms which is monosubstituted or disubstituted by A and/or carbonyl oxygen,
- 35

R³ is H or A,
 Ar is phenyl which is unsubstituted or monosubstituted or
 disubstituted by Hal, A, OR⁴ or O-[C(R⁴)₂]-COOR³,
 5 A is unbranched or branched alkyl having 1-6 carbon atoms, in
 which 1-7 H atoms may be replaced by F,
 n is 0 or 1,
 and their pharmaceutically usable derivatives, solvates and stereo-
 isomers, including mixtures thereof in all ratios.
 10

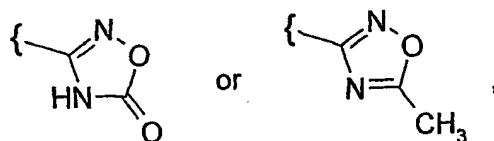
13. Compounds according to Claim 1, in which
 R¹ is -C(=NH)-NH₂ which is unsubstituted or monosubstituted by
 OH, or is
 15  or 
 R² is H,
 20 R³ is H, A or -(CH₂)_n-Ar,
 R⁴ is H or A,
 W is -OC(R³)₂- or -NR³C(R³)₂-,
 X is CONH or CONH(CH₂)₂,
 25 Y is alkylene or Ar-diyl,
 T is dimethylamino, diethylamino, morpholin-4-yl, 2-oxo-
 piperidin-1-yl, 2-oxopyrrolidin-1-yl, 5,5-dimethyl-2-oxo-
 pyrrolidin-1-yl, 2-oxo-1*H*-pyridin-1-yl, 3-oxomorpholin-4-yl, 4-
 30 oxo-1*H*-pyridin-1-yl, 2,6-dioxopiperidin-1-yl, 2-oxopiperazin-1-
 yl, 2,5-dioxopyrrolidin-1-yl, 2-oxo-1,3-oxazolidin-3-yl, 2-aza-
 bicyclo[2.2.2]octan-3-on-2-yl or 3-oxo-2*H*-pyridazin-2-yl,
 Ar is phenyl which is unsubstituted or monosubstituted or
 35 disubstituted by Hal, A, OR⁴ or O-[C(R⁴)₂]-COOR³,

A is unbranched or branched alkyl having 1-6 carbon atoms, in which 1-7 H atoms may be replaced by F,
 n is 0 or 1,
 5 and their pharmaceutically usable derivatives, solvates and stereoisomers, including mixtures thereof in all ratios.

14. Compounds according to Claim 1, in which

R¹ is CN, NH₂, CONA₂, CH₂NH₂, CH₂CH₂NH₂, -C(=NH)-NH₂,
 10 which is unsubstituted or monosubstituted by OH, COOR³, OCOA or OCOOA, or is

15

R²

is H or F,

R³ is H, A or -(CH₂)_n-Ar or thiienyl,R⁴ is H or A,

20

W is -C(R³)₂⁻, -OC(R³)₂⁻ or -NR³C(R³)₂⁻,X is CONH, CONH(CH₂), CONH(CH₂)₂, CH₂O- or CH₂OCH₂,

Y is alkylene, Ar-diyl or pyridinediyl,

25

T is dimethylamino, diethylamino, morpholin-4-yl, 2-oxopiperidin-1-yl, 2-oxopyrrolidin-1-yl, 5,5-dimethyl-2-oxopyrrolidin-1-yl, 2-oxo-1H-pyridin-1-yl, 3-oxomorpholin-4-yl, 4-oxo-1H-pyridin-1-yl, 2,6-dioxopiperidin-1-yl, 2-oxopiperazin-1-yl, 2,5-dioxopyrrolidin-1-yl, 2-oxo-1,3-oxazolidin-3-yl, 3-oxo-2H-pyridazin-2-yl, 2-caprolactam-1-yl, 2-oxo-1,3-oxazinan-3-yl or 2-azabicyclo[2.2.2]octan-3-on-2-yl,

30

Ar is phenyl which is unsubstituted or monosubstituted or disubstituted by Hal, CF₃, A, OA, methoxycarbonylmethoxy, ethoxycarbonylmethoxy or carboxymethoxy,

35

A is unbranched or branched alkyl having 1-6 carbon atoms, in which 1-7 H atoms may be replaced by F,

n is 0 or 1,

5 and their pharmaceutically usable derivatives, solvates and stereoisomers, including mixtures thereof in all ratios.

10 15. Compounds according to Claim 1, selected from the group consisting of

15 2-(3-amidinophenoxy)-N-(4-morpholin-4-ylphenyl)valeramide,

2-(3-amidinophenoxy)-N-(4-morpholin-4-ylphenyl)-2-phenylacetamide,

20 25 2-(3-amidinophenylamino)-N-(4-dimethylaminophenyl)-4-methylvaleramide,

2-(3-amidinophenylamino)-N-(4-morpholin-4-ylphenyl)-2-phenylacetamide,

30 35 2-(3-amidinophenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,

2-(3-amidinophenoxy)-N-[4-(2-oxopyrrolidin-1-yl)phenyl]-2-phenylacetamide;

2-(3-amidinophenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]valeramide,

2-(3-amidinophenoxy)-N-[4-(2-oxopyrrolidin-1-yl)phenyl]valeramide,

2-(3-amidinophenoxy)-N-[4-(2-oxo-1H-pyridin-1-yl)phenyl]-2-phenylacetamide,

2-(3-amidinophenoxy)-N-[4-(3-oxomorpholin-4-yl)phenyl]-2-phenylacetamide,

2-(3-amidinophenoxy)-N-[4-(4-oxo-1H-pyridin-1-yl)phenyl]-2-phenylacetamide,

35 2-(3-(N-hydroxyamidino)phenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,

- 2-(3-amidinophenoxy)-*N*-[4-(2,6-dioxopiperidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-*N*-[4-(3-oxo-2*H*-piperazin-1-yl)phenyl]-2-phenylacetamide,
5 2-(3-amidinophenoxy)-*N*-[2-fluoro-4-(2-oxo-1*H*-pyridin-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-*N*-[4-(2,5-dioxopyrrolidin-1-yl)phenyl]-2-phenylacetamide,
10 2-(3-amidinophenoxy)-*N*-[5-(2-oxopiperidin-1-yl)pyridin-2-yl]-2-phenylacetamide,
2-(3-amidinophenoxy)-*N*-[4-(2-oxo-1,3-oxazolidin-3-yl)phenyl]-2-phenylacetamide,
15 2-(3-amidinophenoxy)-*N*-[4-(2-oxo-1*H*-pyridin-1-yl)benzyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-*N*-[4-(2-oxopiperidin-1-yl)benzyl]-2-phenylacetamide,
20 2-(3-amidinophenoxy)-*N*-[4-(2-oxopiperidin-1-yl)butyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-*N*-[4-(3-oxo-2*H*-pyridazin-2-yl)phenyl]-2-phenylacetamide,
25 2-(3-amidinophenoxy)-*N*-[2-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-*N*-(4-dimethylaminobenzyl)-2-phenylacetamide,
30 2-(3-amidinophenoxy)-*N*-[3-(morpholin-4-yl)propyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-*N*-[3-(piperidin-1-yl)propyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-*N*-(4-dimethylaminophenyl)-2-phenylacetamide,
35

- 2-(3-amidinophenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenylamino)-*N*-[2-(morpholin-4-yl)ethyl]-2-phenylacetamide,
5 2-(3-amidinophenylamino)-*N*-(4-dimethylaminobenzyl)-2-phenylacetamide,
2-(3-amidinophenylamino)-*N*-[3-(morpholin-4-yl)propyl]-2-phenylacetamide,
10 2-(3-amidinophenylamino)-*N*-[3-(piperidin-1-yl)propyl]-2-phenylacetamide
2-(3-amidinophenoxy)-*N*-[2-(morpholin-4-yl)ethyl]valeramide,
2-(3-amidinophenoxy)-*N*-(4-dimethylaminobenzyl)valeramide,
15 2-(3-amidinophenoxy)-*N*-[3-(morpholin-4-yl)propyl]valeramide,
2-(3-amidinophenoxy)-*N*-[3-(piperidin-1-yl)propyl]valeramide,
2-(3-aminocarbonylphenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
20 2-(3-aminocarbonylphenoxy)-*N*-[2-(morpholin-4-yl)ethyl]-2-phenylacetamide,
2-(3-aminocarbonylphenoxy)-*N*-(4-dimethylaminobenzyl)-2-phenylacetamide,
25 2-(3-aminocarbonylphenoxy)-*N*-[3-(morpholin-4-yl)propyl]-2-phenylacetamide,
2-(3-aminocarbonylphenoxy)-*N*-[3-(piperidin-1-yl)propyl]-2-phenylacetamide,
30 2-(3-amidinophenoxy)-*N*-(4-morpholin-4-ylbenzyl)valeramide,
2-(3-amidinophenoxy)-*N*-(4-morpholin-4-ylbenzyl)-2-phenylacetamide,
35 2-(3-amidinophenoxy)-*N*-[4-(3-oxomorpholin-4-yl)phenyl]-2-phenylacetamide,

- 2-(3-amidinophenoxy)-*N*-[3-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
5 (2*R*)-2-(3-amidinophenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]valeramide,
2-(3-amidinophenoxy)-*N*-[2-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
10 2-(3-amidinophenoxy)-*N*-[2-fluoro-4-(2-caprolactam-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-*N*-[3-methyl-4-(2-caprolactam-1-yl)phenyl]-valeramide,
15 2-(3-amidinophenoxy)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-valeramide,
2-(3-amidinophenoxy)-*N*-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-valeramide,
20 2-(3-amidinophenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]acetamide,
2-(2-amidinophenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
25 2-(4-amidinophenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
3-(3-amidinophenyl)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]propionamide,
30 2-(3-amidinophenoxy)-*N*-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
3-(3-amidinophenyl)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-propionamide,
35 (2*S*)-2-(3-amidinophenoxy)-*N*-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-valeramide,
2-(3-amidinophenoxy)-*N*-[3-methyl-4-(2-caprolactam-1-yl)phenyl]-2-phenylacetamide,

- 2-(3-amidinophenylmethyl)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]valeramide,
2-(3-amidinophenylmethyl)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)-
phenyl]valeramide,
5 2-(3-amidinophenoxy)-*N*-[4-(2-oxopiperazin-1-yl)phenyl]valeramide,
(2*S*)-2-(3-amidinophenoxy)-*N*-[4-(2-caprolactam-1-yl)phenyl]valeramide,
10 (2*S*)-2-(3-amidinophenoxy)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)-
phenyl]valeramide,
2-(3-amidinophenoxy)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-4-
15 methylvaleramide,
2-(3-amidinophenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-4-methyl-
valeramide,
(2*R*)-2-(3-amidinophenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]valeramide,
20 2-(3-amidinophenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]propionamide,
2-(3-amidinophenoxy)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-
propionamide,
2-(3-amidinophenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]butyramide,
2-(3-amidinophenoxy)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-
25 butyramide,
(2*S*)-2-(3-amidinophenoxy)-*N*-[3-methyl-4-(2-caprolactam-1-yl)-
phenyl]valeramide,
2-(3-amidinophenoxy)-*N*-[4-(2-oxopyridin-1-yl)phenyl]valeramide,
30 2-(3-amidinophenoxy)-*N*-[2,5-dimethyl-4-(2-oxopiperidin-1-yl)phenyl]-
valeramide,
2-(3-amidinophenoxy)-*N*-[3-methyl-4-(2-oxopyrrolidin-1-yl)phenyl]-2-
phenylacetamide,
35 2-(3-amidinophenoxy)-*N*-[3-methyl-4-(2-oxopyrrolidin-1-yl)phenyl]-
valeramide,

- 2-(3-amidinophenoxy)-*N*-[2-methyl-4-(2-oxopiperidin-1-yl)phenyl]-valeramide,
2-(3-amidinophenoxy)-*N*-(2-oxo-3,4,5,6-tetrahydro-2*H*-1,2'-bi-pyridinyl-5'-yl)valeramide,
5 2-(3-amidinophenoxy)-*N*-[2-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-*N*-(2-oxo-3,4,5,6-tetrahydro-2*H*-1,2'-bi-pyridinyl-5'-yl)-2-phenylacetamide,
10 2-[*N*-(3-*N*-methoxycarbonylamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]valeramide,
2-(3-amidinophenoxy)-*N*-[3-fluoro-4-(2-oxopyrrolidin-1-yl)phenyl]valeramide,
15 2-(3-amidinophenoxy)-*N*-[3-fluoro-4-(2-oxopyrrolidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]caproamide,
2-(3-amidinophenoxy)-*N*-[4-(2-oxopyrrolidin-1-yl)phenyl]butyramide,
20 2-(3-amidinophenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-3-methylbutyramide,
2-(3-amidinophenoxy)-*N*-[4-ethyl-3-(2-oxopyrrolidin-1-yl)phenyl]valeramide,
25 2-(3-amidinophenoxy)-*N*-[3-fluoro-4-(2-caprolactam-1-yl)phenyl]valeramide,
2-(3-amidinophenoxy)-*N*-[3-fluoro-4-(2-caprolactam-1-yl)phenyl]-2-phenylacetamide, acetate,
30 2-(3-amidinophenoxy)-*N*-[4-(2,6-dioxopiperidin-1-yl)phenyl]valeramide,
3-(3-amidinophenyl)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylpropionamide,
35 3-(3-amidinophenyl)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylpropionamide,

- 2-(3-amidinophenoxy)-*N*-[3-methoxy-4-(2,5-dioxopyrrolidin-1-yl)-phenyl]valeramide,
1-[4-[2-(3-amidinophenoxy)pentylamino]phenyl]piperidin-2-one,
1-[4-[2-(3-amidinophenoxy)pentylamino]-2-methylphenyl]piperidin-2-one,
1-[4-[2-(3-amidinophenoxy)-2-phenylethoxy]phenyl]piperidin-2-one,
2-(3-amidinophenoxy)-*N*-[3-(2-oxopiperidin-1-yl)propyl]-2-phenylacetamide,
2-[(3-*N*-ethoxycarbonylamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]valeramide,
2-[(3-*N*-methoxycarbonylamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
2-[(3-*N*-ethoxycarbonylamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide,
2-(3-amidinophenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenylamino)-*N*-[4-(2-oxopyrrolidin-1-yl)phenyl]-4-methylvaleramide,
2-(3-amidinophenylamino)-*N*-[4-(2-oxopyrrolidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
2-(3-amidinophenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]acetamide,
(2*S*)-2-(3-amidinophenylamino)-*N*-[4-(2-caprolactam-1-yl)phenyl]-2-phenylacetamide,
(2*S*)-2-(3-amidinophenylamino)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,

- (2S)-2-(3-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]caproamide,
5 2-(3-amidinophenylamino)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
(2S)-2-(2-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
10 (2R)-2-(2-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
(2S)-2-(3-amidinophenylamino)-N-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
15 2-(3-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-(3-fluorophenyl)acetamide,
2-(3-amidinophenylamino)-N-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-2-(3-fluorophenyl)acetamide,
20 (2R)-2-(3-amidinophenylamino)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide,
2-(3-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]valeramide,
25 (2R)-2-(3-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide,
(2R)-2-(3-amidinophenylamino)-N-[4-(2-caprolactam-1-yl)phenyl]-4-methylvaleramide,
30 2-(3-amidinophenylamino)-N-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
2-(3-amidinophenylamino)-N-[3-methyl-4-(2-caprolactam-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
35 (2R)-2-(3-amidinophenylamino)-N-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide,

- 2-(3-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-4-trifluoro-3-methylbutyramide,
5 (2S)-2-(3-amidinophenylamino)-N-[4-(2-oxopiperidin-1-yl)benzyl]-2-phenylacetamide,
(2S)-2-(3-amidinophenylamino)-N-[4-(3-oxo-2-azabicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide,
10 2-(3-amidinophenylamino)-N-[3-trifluoromethyl-4-(3-oxo-2-azabicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenylamino)-N-methyl-N-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide,
15 2-(3-aminocarbonylphenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]valeramide,
2-(3-aminocarbonylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
20 2-(3-aminocarbonylphenoxy)-N-[4-(2-oxo-1H-pyridin-1-yl)benzyl]-2-phenylacetamide,
2-(3-aminocarbonylphenoxy)-N-[4-(2-caprolactam-1-yl)phenyl]-2-phenylacetamide,
25 2-(3-aminocarbonylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
3-(3-aminocarbonylphenyl)-N-[4-(2-oxopiperidin-1-yl)phenyl]propionamide,
30 2-(3-aminocarbonylphenoxy)-N-[4-(2-oxo-1H-pyridin-1-yl)phenyl]-2-phenylacetamide,
2-(3-aminocarbonylphenoxy)-N-[3-methyl-4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide,
35 2-(3-aminocarbonylphenoxy)-N-[4-(4-oxo-4H-pyridin-1-yl)phenyl]-2-phenylacetamide,

- 2-(3-aminocarbonylphenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-acetamide,
5 (2S)-2-(3-aminocarbonylphenylamino)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
(2S)-2-(3-aminocarbonylphenylamino)-*N*-[4-(2-caprolactam-1-yl)-phenyl]-2-phenylacetamide,
10 (2R)-2-(3-aminocarbonylphenylamino)-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide,
(2S)-2-(3-aminocarbonylphenylamino)-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide,
15 2-(3-aminocarbonylphenylamino)-*N*-[4-(2-caprolactam-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
2-(3-aminocarbonylphenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-(2-thienyl)acetamide,
20 2-(4-aminocarbonylphenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
2-(2-aminocarbonylphenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
25 2-(3-aminocarbonylphenoxy)-*N*-[2-fluoro-4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide,
2-(3-aminocarbonylphenoxy)-*N*-[2-fluoro-4-(2-caprolactam-1-yl)-phenyl]-2-phenylacetamide,
30 (2S)-2-(2-aminocarbonylphenyl)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-aminocarbonylphenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
35 (2S)-2-(3-aminocarbonylphenylamino)-*N*-[4-(2-oxopyridin-1-yl)-phenyl]-2-phenylacetamide,

- 2-(3-aminocarbonylphenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-valeramide,
5 2-(3-aminocarbonyl-4-fluorophenylamino)-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide,
2-(3-aminocarbonylphenylamino)-*N*-[3-trifluoromethyl-4-(3-oxo-2-azabicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide,
10 2-(3-*N,N*-diethylaminocarbonylphenoxy)-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide,
15 2-(3-aminocarbonylphenoxy)-*N*-[4-(2-oxopiperazin-1-yl)phenyl]-2-phenylacetamide,
2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide,
20 2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
2-[3-(*N*-hydroxyamidino)phenyl]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-propionamide,
25 (2*R*)-[2-(3-(*N*-hydroxyamidino)phenylamino)-*N*-[4-(2-caprolactam-1-yl)phenyl]-2-phenylacetamide,
2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]-2-(2-trifluoromethylphenyl)acetamide,
30 2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]-2-(2-thienyl)acetamide,
35 2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)-phenyl]valeramide,

- (2S)-2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[3-fluoro-4-(2-oxo-piperidin-1-yl)phenyl]-2-phenylacetamide,
3-[3-(*N*-hydroxyamidino)phenyl]-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)-phenyl]propionamide,
5 (2S)-2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]-4-methylvaleramide,
2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-valeramide,
10 2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxo-1*H*-pyrazin-1-yl)-phenyl]valeramide,
(2S)-2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide,
15 (2R)-2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide,
2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]valeramide,
20 2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide,
2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxopyridin-1-yl)phenyl]-valeramide,
25 (2R)-2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-caprolactam-1-yl)-phenyl]-4-methylvaleramide,
(2R)-2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]valeramide,
30 (2S)-2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]valeramide,
2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide,
35 2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]-3-methyl-4-trifluorobutyramide,

- (2S)-2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)benzyl]-2-phenylacetamide,
5 (2S)-2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[4-(3-oxo-2-azabicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide,
2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-butyramide,
10 2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-propionamide,
15 2-[3-(*N*-hydroxyamidino)phenylamino]-*N*-[3-trifluoromethyl-4-(3-oxo-2-azabicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide,
3-[3-(*N*-hydroxyamidino)phenyl]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylpropionamide,
20 3-[3-(*N*-hydroxyamidino)phenyl]-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylpropionamide,
1-{4-[2-(3-(*N*-hydroxyamidino)phenoxy)-2-phenylethoxy]phenyl}-piperidin-2-one,
25 2-[3-(*N*-methoxycarbonyloxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]valeramide,
2-[3-(*N*-ethoxycarbonyloxyamidino)phenoxy]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]valeramide,
30 (2S)-2-[3-(*N*-ethoxycarbonyloxyamidino)phenylamino]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide,
2-(3-aminomethylphenylamino)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
35 2-(3-aminomethylphenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, hydrochloride,
2-(3-aminomethylphenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
3-(3-aminomethylphenyl)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]propionamide,

- 2-(3-aminomethylphenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]acetamide,
2-(3-aminomethylphenoxy)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]valeramide,
5 (2*S*)-2-(3-aminomethylphenylamino)-*N*-[4-(2-caprolactam-1-yl)-phenyl]-2-phenylacetamide,
(2*S*)-2-(3-aminomethylphenylamino)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]acetamide,
10 2-(3-aminomethylphenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-(2-trifluoromethylphenyl)acetamide,
(2*S*)-2-(3-aminomethylphenylamino)-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide,
15 (2*R*)-2-(3-aminomethylphenylamino)-*N*-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide,
2-(3-aminomethylphenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-(2-methylphenyl)acetamide,
20 2-(3-aminomethylphenylamino)-*N*-[4-(2-caprolactam-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
2-(3-aminomethylphenylamino)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)-phenyl]-2-(2-methylphenyl)acetamide,
25 2-(3-aminomethylphenylamino)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide,
2-(3-aminomethylphenoxy)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)-phenyl]valeramide,
30 2-(3-aminomethylphenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-(2-thienyl)acetamide,
2-(3-aminomethylphenylamino)-*N*-[3-fluoro-4-(2-oxopiperidin-1-yl)-phenyl]-2-(3-fluorophenyl)acetamide,
35 (2*S*)-2-(3-aminomethylphenylamino)-*N*-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,

- (2S)-2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
5
(2S)-2-(2-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide,
2-(2-aminomethylphenyl)-N-[4-(2-oxopiperidin-1-yl)phenyl]acetamide,
2-(2-aminomethylphenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
10
2-(4-aminomethylphenoxy)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-(3-fluorophenyl)acetamide,
15
2-(3-aminomethylphenoxy)-N-[3-methyl-4-(2-caprolactam-1-yl)-phenyl]-2-phenylacetamide,
(2R)-2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-caprolactam-1-yl)phenyl]-2-phenylacetamide,
20
(2R)-2-(2-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide,
(2S)-2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-caprolactam-1-yl)phenyl]-2-phenylacetamide,
25
2-(3-aminomethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-cyclohexylacetamide,
2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-oxopiperidin-1-yl)-phenyl]-2-cyclohexylacetamide,
30
2-(3-aminomethylphenylamino)-N-[4-(2-caprolactam-1-yl)phenyl]-4-methylvaleramide,
(2R)-2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, hydrochloride,
(2S)-2-(3-aminomethylphenylamino)-N-[4-(2-oxopyridin-1-yl)phenyl]-2-phenylacetamide,
35

- 2-(3-aminomethylphenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-valeramide,
2-(3-aminomethylphenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide,
5 (2*R*)-2-(3-aminomethylphenylamino)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide,
(2*R*)-2-(3-aminomethylphenylamino)-*N*-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide,
10 (2*R*)-2-(3-aminomethylphenylamino)-*N*-[4-(2-caprolactam-1-yl)phenyl]-4-methylvaleramide,
(2*S*)-2-(3-aminomethylphenylamino)-*N*-[2,5-dimethyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
15 2-(3-aminomethylphenylamino)-*N*-[3-fluoro-4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
2-(3-aminomethylphenylamino)-*N*-[3-methyl-4-(2-caprolactam-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
20 2-(3-aminomethylphenylamino)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-(2,4-difluorophenyl)acetamide,
2-(3-aminomethylphenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-(2,4-difluorophenyl)acetamide,
25 (2*S*)-2-(3-aminomethylphenylamino)-*N*-[3-methyl-4-(2-oxopyrrolidin-1-yl)phenyl]-2-phenylacetamide,
(2*S*)-2-(3-aminomethylphenylamino)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-4-methylvaleramide,
30 2-(3-aminomethylphenylamino)-*N*-methyl-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-aminoethylphenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide, hydrochloride,
35 2-(3-aminomethylphenylamino)-*N*-[3-methyl-4-(2-oxopyrrolidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,

- 2-(3-aminomethylphenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-4-trifluoro-3-methylbutyramide,
2-(3-aminomethylphenylamino)-*N*-[4-(2-oxopiperidin-1-yl)benzyl]-2-phenylacetamide,
5 (2*S*)-2-(3-aminomethylphenylamino)-*N*-[4-(3-oxo-2-azabicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide,
2-(3-aminomethylphenylamino)-*N*-[3-fluoro-4-(2-oxopyrrolidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
10 2-(3-aminomethylphenylamino)-*N*-[4-(2-oxopyrrolidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-aminomethylphenylamino)-*N*-[4-(2-oxopyrrolidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
15 2-(3-aminomethylphenylamino)-*N*-[4-ethyl-3-(2-oxopyrrolidin-1-yl)phenyl]-2-phenylacetamide,
(2*S*)-2-(3-aminomethylphenylamino)-*N*-[3-methoxy-4-(3-oxo-2-aza-
bicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide,
20 2-(3-aminomethylphenylamino)-*N*-[3-fluoro-4-(2-caprolactam-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
(2*S*)-2-(3-aminomethylphenylamino)-*N*-[3-fluoro-4-(2-caprolactam-1-
yl)phenyl]-2-phenylacetamide,
25 (2*S*)-2-(3-aminomethylphenylamino)-*N*-[3-trifluoromethyl-4-(3-oxo-2-
azabicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide,
2-(3-aminomethylphenylamino)-*N*-[3-methyl-4-(3-oxo-2-azabicyclo-
[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide,
30 (2*S*)-2-(3-aminomethylphenylamino)-*N*-[3-methoxy-4-(3-oxo-2-aza-
bicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide,
2-(3-aminomethylphenylamino)-*N*-[3-trifluoromethyl-4-(2-oxopiperidin-
1-yl)phenyl]-2-phenylacetamide,
35 2-(3-aminomethylphenylamino)-*N*-[3-methoxy-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,

- 2-(3-aminomethylphenylamino)-N-[3-methyl-4-(3-oxo-2-azabicyclo-[2.2.2]oct-2-yl)phenyl]-2-(2-fluorophenyl)acetamide,
5
2-(3-aminomethyl-4-fluorophenylamino)-N-[4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide,
2-(3-aminomethylphenylamino)-N-[3-methyl-4-(3-oxo-2-azabicyclo-[2.2.2]oct-2-yl)phenyl]-2-(2-fluorophenyl)acetamide,
10
2-(3-aminomethylphenylamino)-N-[3-chloro-4-(2-oxopyrrolidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide,
15
2-(3-aminomethylphenylamino)-N-[3-methoxy-4-(2-oxopiperidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide,
2-(3-aminomethylphenylamino)-N-[3-trifluoromethyl-4-(3-oxo-2-azabicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide,
20
2-(3-aminomethylphenylamino)-N-[3-trifluoromethyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
3-(3-aminomethylphenyl)-N-[4-(2-caprolactam-1-yl)phenyl]-2-phenylpropionamide,
25
2-(3-aminomethylphenoxy)-N-[3-methyl-4-(2-oxopyrrolidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide,
3-(3-aminomethylphenyl)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylpropionamide,
30
2-(3-aminomethylphenoxy)-N-[3-methyl-4-(2-caprolactam-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide,
2-(3-aminomethylphenoxy)-N-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
35
3-(3-aminomethylphenyl)-N-[3-methyl-4-(2-oxopyrrolidin-1-yl)phenyl]-2-phenylpropionamide,

- (2S)-2-(3-aminomethylphenylamino)-N-[4-(5,5-dimethyl-2-oxo-pyrrolidin-1-yl)phenyl]-2-phenylacetamide,
5 (2S)-2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-oxo-1,3-oxazinan-3-yl)phenyl]-2-phenylacetamide,
(2S)-2-(3-aminomethylphenylamino)-N-[3-methyl-4-(2-oxooxazolidin-3-yl)phenyl]-2-phenylacetamide,
10 (2S)-2-(3-aminoethylphenylamino)-N-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-N-[4-(2-oxopiperazin-1-yl)phenyl]-2-phenylacetamide,
15 2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]-N-[4-(2,6-dioxopiperidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-N-[4-(2,6-dioxopiperidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-N-[3-trifluoromethyl-4-(2,6-dioxopiperidin-1-yl)phenyl]valeramide,
20 2-(3-amidinophenoxy)-N-[3-chloro-4-(2,5-dioxopyrrolidin-1-yl)phenyl]valeramide, acetate,
2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]-N-[3-trifluoromethyl-4-(2,5-dioxopyrrolidin-1-yl)phenyl]-2-valeramide,
25 2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]-N-[3-chloro-4-(2,5-dioxopyrrolidin-1-yl)phenyl]-2-valeramide,
2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]-N-[3-methoxy-4-(2,5-dioxopyrrolidin-1-yl)phenyl]-2-valeramide,
30 2-[3-(5-methyl-1,2,4-oxadiazol-3-yl)phenoxy]-N-[4-(2,5-dioxopyrrolidin-1-yl)phenyl]-2-valeramide,
3-(3-amidinophenoxy)-N-[3-chloro-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
35 3-(3-amidinophenoxy)-N-[3-chloro-4-(2-oxopiperidin-1-yl)phenyl]valeramide,

- 3-(3-amidinophenoxy)-*N*-[3-chloro-4-(2-oxopiperidin-1-yl)phenyl]-butyramide,
2-(3-amidino-4-fluorophenylamino)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
5 3-(3-aminomethylphenyl)-*N*-[3-methyl-4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)propionamide,
3-(3-aminomethylphenyl)-*N*-[4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)propionamide,
10 3-(3-aminomethylphenylamino)-*N*-[3-chloro-4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
3-(3-aminomethylphenylamino)-*N*-[3-chloro-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
15 3-(3-aminomethylphenoxy)-*N*-[3-chloro-4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
3-(3-aminomethylphenyl)-*N*-[3-methyl-4-(2-oxopyrrolidin-1-yl)phenyl]-2-(2-fluorophenyl)propionamide,
20 3-(3-aminomethylphenyl)-*N*-[3-chloro-4-(2-oxopiperidin-1-yl)phenyl]-2-(2-fluorophenyl)propionamide,
3-(3-aminomethylphenylamino)-*N*-[4-(2,6-dioxopiperidin-1-yl)phenyl]-2-phenylacetamide, trifluoroacetate, ESI 443;
25 2-(3-aminomethylphenylamino)-*N*-[3-chloro-4-(2,5-dioxopyrrolidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-aminomethyl-4-fluorophenylamino)-*N*-[3-methyl-4-(3-oxo-2-aza-bicyclo[2.2.2]oct-2-yl)phenyl]-2-phenylacetamide,
30 2-[3-(*N*-tertbutoxycarbonylaminomethyl)phenylamino]-*N*-[3-chloro-4-(2,5-dioxopyrrolidin-1-yl)phenyl]-2-phenylacetamide,
2-(3-amidinophenoxy)-*N*-[3-carboxymethoxy-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
35 2-(3-amidinophenoxy)-*N*-[3-ethoxycarbonylmethoxy-4-(2-oxo-piperidin-1-yl)phenyl]-2-phenylacetamide,

- 2-(3-amidinophenoxy)-*N*-[3-ethoxycarbonylmethoxy-4-(2-oxo-piperidin-1-yl)phenyl]valeramide,
3-(3-amidinophenoxy)-*N*-[3-carboxymethoxy-4-(2-oxopiperidin-1-yl)-phenyl]valeramide,
5 3-(3-amidinophenoxy)-*N*-[3-carboxymethoxy-4-(2-oxopiperidin-1-yl)-phenyl]-2-phenylacetamide,
2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[3-ethoxycarbonylmethoxy-4-(2-oxopiperidin-1-yl)phenyl]-2-phenylacetamide,
10 2-[3-(*N*-hydroxyamidino)phenoxy]-*N*-[3-ethoxycarbonylmethoxy-4-(2-oxopiperidin-1-yl)phenyl]valeramide,
2-(3-aminomethylphenylamino)-*N*-[3-methyl-4-(3-oxo-2-azabicyclo[2.2.2]oct-2-yl)phenyl]-2-(2-fluorophenyl)acetamide,
15 2-(3-aminomethylphenylamino)-*N*-[3-methyl-4-(2-oxooxazolidin-3-yl)-phenyl]-2-(2-fluorophenyl)acetamide,
3-(3-aminomethylphenylamino)-*N*-[3-methoxy-4-(2-oxopiperidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide,
20 3-(3-aminomethylphenylamino)-*N*-[3-chloro-4-(2-oxopyrrolidin-1-yl)-phenyl]-2-phenylacetamide,
2-(3-aminomethylphenoxy)-*N*-[3-methyl-4-(3-oxo-2-azabicyclo[2.2.2]oct-2-yl)phenyl]-2-(2-fluorophenyl)acetamide,
25 3-(3-aminomethylphenylamino)-*N*-[4-(5,5-dimethyl-2-oxopyrrolidin-1-yl)phenyl]-2-(2-fluorophenyl)acetamide,
3-(3-aminomethylphenoxy)-*N*-[4-(5,5-dimethyl-2-oxopyrrolidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide,
30 2-(3-aminomethylphenoxy)-*N*-[3-methyl-4-(2-oxooxazolidin-3-yl)-phenyl]-2-(2-fluorophenyl)acetamide,
3-(3-aminomethylphenoxy)-*N*-[3-methoxy-4-(2-oxopiperidin-1-yl)-phenyl]-2-(2-fluorophenyl)acetamide,

and their pharmaceutically usable derivatives, solvates and stereoisomers, including mixtures thereof in all ratios.

- 5 16. Process for the preparation of compounds of the formula I according
to Claim 1 and their pharmaceutically tolerated salts and solvates,
characterised in that
- 10 a) they are liberated from one of their functional derivatives by
treatment with a solvolysing or hydrogenolysing agent by
- 15 i) liberating an amidino group from their hydroxyl, oxadiazole or
oxazolidinone derivative by hydrogenolysis or solvolysis,
- 20 ii) replacing a conventional amino-protecting group by hydrogen
by treatment with a solvolysing or hydrogenolysing agent, or
liberating an amino group protected by a conventional
protecting group,

or

- 25 b) converting a cyano group into an amidino group,

or

- 30 c) reacting a compound of the formula II



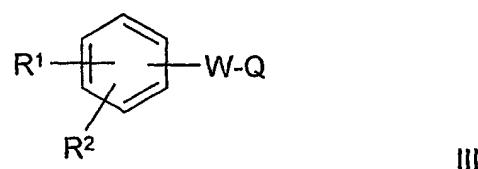
35 in which

Z is HNR^3- or $\text{HNR}^3\text{C}(\text{R}^4)_2-$

and R³, R⁴, Y and T are as defined in Claim 1,

with a compound of the formula III

5

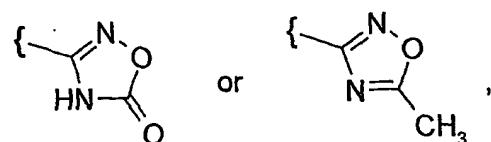


10

in which

15

R¹ is -C(=NH)-NH₂ which is monosubstituted by C(=O)R³, COOR³, OR³ or by a conventional amino-protecting group, or is



20

Q is -CO-L or -C(R⁴)₂-L,

L is Cl, Br, I or a free or reactively functionally modified OH group, and

25

R², R³, R⁴ and W are as defined in Claim 1,

and/or

30

d) converting a base or acid of the formula I into one of its salts.

35

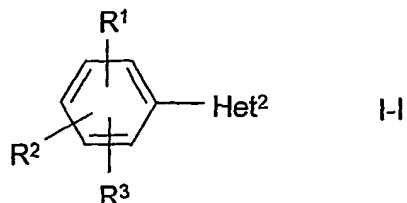
17. Compounds of the formula I according to one or more of Claims 1 to 15 as inhibitors of coagulation factor Xa.

18. Compounds of the formula I according to one or more of Claims 1 to 15 as inhibitors of coagulation factor VIIa.
- 5 19. Medicament comprising at least one compound of the formula I according to one or more of Claims 1 to 15 and/or its pharmaceutically usable derivatives, solvates and stereoisomers, including mixtures thereof in all ratios, and, if desired, excipients and/or assistants.
- 10 20. Medicament comprising at least one compound of the formula I according to one or more of Claims 1 to 15 and/or its pharmaceutically usable derivatives, solvates and stereoisomers, including mixtures thereof in all ratios, and at least one further medicament active ingredient.
- 15 21. Use of compounds according to one or more of Claims 1 to 15 and/or their physiologically acceptable salts and solvates for the preparation of a medicament for the treatment of thrombosis, myocardial infarction, arteriosclerosis, inflammation, apoplexia, angina pectoris, restenosis after angioplasty, claudicatio intermittens, tumours, tumour diseases and/or tumour metastases.
- 20 22. Set (kit) consisting of separate packs of
 - (a) an effective amount of a compound of the formula I according to one or more of Claims 1 to 15 and/or its pharmaceutically usable derivatives, solvates and stereoisomers, including mixtures thereof in all ratios,
- 25 and
 - (b) an effective amount of a further medicament active ingredient.
- 30 35

23. Use of compounds of the formula I according to one or more of
 5 Claims 1 to 15 and/or their pharmaceutically usable derivatives,
 solvates and stereoisomers, including mixtures thereof in all ratios,
 for the preparation of a medicament for the treatment of thrombosis,
 myocardial infarction, arteriosclerosis, inflammation, apoplexia,
 angina pectoris, restenosis after angioplasty, claudicatio intermittens,
 tumours, tumour diseases and/or tumour metastases,
 10 in combination with at least one further medicament active ingredient.
 VI

24. Intermediates of the formula I-I

15



20

in which

25

R¹ is NO₂, CN, NHA, NHCOA, NACF₃, NH₂ or (CH₂)_nNH₂,

R² and

R³ are each, independently of one another, H, Hal, A, OA or CF₃,

30

A is alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,

Het² is a monocyclic or bicyclic, saturated, unsaturated or aromatic heterocyclic radical having from 1 to 4 N, O and/or S atoms

which is unsubstituted or monosubstituted, disubstituted or trisubstituted by Hal, A, OA, CN, COOA, CONH₂, NHCOA,

NHSO₂A, S(O)_mA and/or carbonyl oxygen,

n is 0, 1, 2 or 3,

35

and salts thereof.

25. Compounds according to Claim 24, selected from the group consisting of

- 5 1-(3-fluoro-4-nitrophenyl)piperidin-2-one,
 1-(4-amino-3-fluorophenyl)piperidin-2-one,
 tert-butyl 4-(4-nitrophenyl)-3-oxopiperazin-1-carboxylate,
 tert-butyl 4-(4-aminophenyl)-3-oxopiperazin-1-carboxylate,
 1-(4-nitrophenyl)azepam-2-one,
10 1-(4-aminophenyl)azepam-2-one,
 1-(3-fluoro-4-nitrophenyl)azepam-2-one,
 1-(4-amino-3-fluorophenyl)azepam-2-one,
 1-(2-methyl-4-nitrophenyl)azepam-2-one,
15 1-(4-amino-2-methylphenyl)azepam-2-one,
 1-(4-nitrophenyl)-1*H*-pyrazin-2-one,
 1-(4-aminophenyl)-1*H*-pyrazin-2-one,
 1-(2,5-dimethyl-4-nitrophenyl)piperidin-2-one,
20 1-(4-amino-2,5-dimethylphenyl)piperidin-2-one,
 2,2,2-trifluoro-N-methyl-N-[4-(2-oxopiperidin-1-yl)phenyl]acetamide,
 1-(4-methylaminophenyl)piperidin-2-one,
 1-(3-methyl-4-nitrophenyl)piperidin-2-one,
25 1-(4-amino-3-methylphenyl)piperidin-2-one,
 5'-nitro-3,4,5,6-tetrahydro-1,2'-bipyridinyl-2-one,
 5'-amino-3,4,5,6-tetrahydro-1,2'-bipyridinyl-2-one,
 2-(4-nitrophenyl)-2-azabicyclo[2.2.2]octan-3-one,
30 2-(4-aminophenyl)-2-azabicyclo[2.2.2]octan-3-one,
 2-(4-nitro-2-trifluoromethylphenyl)-2-azabicyclo[2.2.2]octan-3-one,
 2-(4-amino-2-trifluoromethylphenyl)-2-azabicyclo[2.2.2]octan-3-one,
 2-(2-methyl-4-nitrophenyl)-2-azabicyclo[2.2.2]octan-3-one,
 2-(4-amino-2-methylphenyl)-2-azabicyclo[2.2.2]octan-3-one,
35 2-(2-methoxy-4-nitrophenyl)-2-azabicyclo[2.2.2]octan-3-one,

2-(4-amino-2-methoxyphenyl)-2-azabicyclo[2.2.2]octan-3-one,
1-(2-methoxy-4-nitrophenyl)piperidin-2-one,
1-(4-amino-2-methoxyphenyl)piperidin-2-one,
1-(4-nitro-2-trifluoromethylphenyl)piperidin-2-one,
5 1-(4-amino-2-trifluoromethylphenyl)piperidin-2-one,
3-(2-methyl-4-nitrophenyl)-1,3-oxazinan-2-one,
3-(4-amino-2-methylphenyl)-1,3-oxazinan-2-one,
1-(2-chloro-4-nitrophenyl)pyrrolidin-2-one,
10 1-(4-amino-2-chlorophenyl)pyrrolidin-2-one,
1-(2-methoxy-4-nitrophenyl)pyrrolidine-2,5-dione,
1-(4-amino-2-methoxyphenyl)pyrrolidine-2,5-dione,
5,5-dimethyl-1-(4-nitrophenyl)pyrrolidin-2-one,
15 1-(4-aminophenyl)-5,5-dimethylpyrrolidin-2-one,
3-(2-methyl-4-nitrophenyl)oxazolidin-2-one,
3-(4-amino-2-methylphenyl)oxazolidin-2-one,
1-(2-fluoro-4-nitrophenyl)azepam-2-one,
20 1-(4-amino-2-fluorophenyl)azepam-2-one,
4-(2-oxo-2H-pyridin-1-yl)benzonitrile,
1-(4-aminomethylphenyl)-1H-pyridin-2-one,
1-(4-aminomethylphenyl)piperidin-2-one,
25 1-(2-ethyl-5-nitrophenyl)pyrrolidin-2-one,
1-(5-amino-2-ethylphenyl)pyrrolidin-2-one,
1-(4-aminophenyl)piperidine-2,6-dione,
and salts thereof.

INTERNATIONAL SEARCH REPORT

Internal Application No
PCT/ [REDACTED] 1/14296

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07D211/76 C07D211/74 C07D265/32 C07D241/08 C07D401/04
 C07D263/22 C07D237/14 C07D223/10 A61K31/535 A61K31/50
 A61K31/4412 A61K31/421

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, CHEM ABS Data, BEILSTEIN Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	TING SU ET AL: "Design and Synthesis of Glycolic and Mandelic Acid derivatives as Factor Xa Inhibitors" BIOORGANIC AND MEDICINAL CHEMISTRY LETTERS, vol. 11, no. 17, 2001, pages 2279-2282, XP001069180 see compounds 19-22	1-23
P,Y	---	1-23
Y	WO 00 71510 A (COR THERAPEUTICS INC) 30 November 2000 (2000-11-30)	1-23
X	see general formula, definitions of A and pages 39/40 and 49 of the description, and example 25 and claims 9,7,6,5. --- -/-	1-23

 Further documents are listed in the continuation of box C. Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "&" document member of the same patent family

Date of the actual completion of the International search

20 June 2002

Date of mailing of the International search report

01.07.02

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
 NL - 2280 HV Rijswijk
 Tel. (+31-70) 340-2040, Tx. 31 651 epo nl
 Fax: (+31-70) 340-3016

Authorized officer

Scruton-Evans, I

INTERNATIONAL SEARCH REPORT

Intern'l Application No
PCT/ [REDACTED] 1/14296

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
E	WO 02 06269 A (MERCK PATENT GMBH ;DORSCH DIETER (DE); GLEITZ JOHANNES (DE); JURAS) 24 January 2002 (2002-01-24) see especially definitions of R5 -----	1-23
A	WO 99 16751 A (BERNOTAT DANIELOWSKI SABINE ;MERCK PATENT GMBH (DE); DORSCH DIETER) 8 April 1999 (1999-04-08) see the definition for R3 and X -----	1-23

INTERNATIONAL SEARCH REPORT

International application No.
PCT/EP 01/14296

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:

2. Claims Nos.: 1-14, 16-23 (partly), 24 because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
see FURTHER INFORMATION sheet PCT/ISA/210

3. Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

As a result of the prior review under R. 40.2(e) PCT,
no additional fees are to be refunded.

1. As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.

2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:

4. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

The additional search fees were accompanied by the applicant's protest.

No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: 1-14,16-23 (partly),24

The initial phase of the search revealed a very large number of documents relevant to the issue of novelty. So many documents were retrieved that it is impossible to determine which parts of the claim(s) may be said to define subject-matter for which protection might legitimately be sought (Article 6 PCT). For these reasons, a meaningful search over the whole breadth of the claim(s) is impossible. Consequently, the search has been restricted to those compounds for which support within the meaning of Article 6 PCT and/or disclosure within the meaning of Article 5 PCT is to be found. The search has thus been carried out for those parts of the claims which appear to be supported and disclosed, namely those parts relating to the compounds wherein R1 is in the 3-position, the linker chain W-X-Y is

-W-C(phenyl or n-propyl)-CON-phenyl, 4-substituted by an N atom (possibly cyclic) and the examples.

For claim 24 the initial phase of the search revealed a very large number of documents relevant to the issue of novelty. So many documents were retrieved that it is impossible to determine which parts of the claim(s) may be said to define subject-matter for which protection might legitimately be sought (Article 6 PCT). For these reasons, a meaningful search over the whole breadth of the claim is impossible. Many known compounds are included within the very general formula, and even making a restriction to a generalisation over the examples (i.e. having a N-bonded 2-carbonyl substituted group for Het, with NO₂ in the 4-position of the phenyl) revealed a very large number of documents relevant to novelty. Thus only the specific examples of claim 25 were searched.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. Claims: 1-23

Compounds of the formula I, their preparation and their use

2. Claims: 24,25

Intermediates of the formula I-I

INTERNATIONAL SEARCH REPORT

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